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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:01; Search time 36.7714 Seconds
(without alignments)
112.231 Million cell updates/sec

Title: US-09-544-664-1

Perfect score: 138

Sequence: 1 NLMAAQRGRELRRMSDEPGFKGL (26)

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: A_Geneseq_19Jun03.*

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21: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2000.DAT.*
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23: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2002.DAT.*
24: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	138	100.0	26	21 AAB37001	Bcl2 polypeptide B
2	138	100.0	26	21 AAB37002	Bcl2 polypeptide B
3	138	100.0	27	21 AAB37003	Bcl2 polypeptide B
4	138	100.0	27	21 AAB37056	Bcl2 polypeptide B
5	138	100.0	28	21 AAB37055	Bcl2 polypeptide B
6	138	100.0	162	22 AAB70370	Shorter murine BAD
7	138	100.0	204	17 AAR95168	bcl-x(L)/bcl-2 ass
8	138	100.0	204	17 AAW61315	Murine BCL-XL/BCL-
9	138	100.0	204	19 AAW61316	Mutant BCL-XL/BCL-

10	138	100.0	204	19 AAW61317	Mutant BCL-XL/BCL-
11	138	100.0	204	19 AAW61318	Mutant BCL-XL/BCL-
12	138	100.0	204	19 AAW58832	Murine BAD protein
13	138	100.0	204	22 AAB70369	Longer murine BAD
14	138	100.0	204	24 ABR39082	Murine BAD protein
15	138	100.0	204	24 ABR39082	Bad-DTRR apoptosis
16	114	82.6	24	22 AAU00220	Human Bad peptide
17	114	82.6	25	23 ABP56161	PTPC-interacting T
18	114	82.6	25	23 ABG78484	Mutant Bcl2 compet
19	114	82.6	25	23 ABG78493	Mutant Bcl2 compet
20	114	82.6	25	23 AAU78610	Human Bad peptide
21	114	82.6	25	23 AAU78620	BBC6 protein for r
22	114	82.6	166	18 AAW32476	Human Bcl-XL/Bcl-2
23	114	82.6	168	19 AAW57779	Human cell prolif
24	114	82.6	168	21 AAB13512	Human BAD mutant a
25	114	82.6	168	22 AAB70368	Human BAD protein.
26	114	82.6	168	22 AAB48287	Amino acid sequenc
27	114	82.6	168	22 AAG67688	Human BAD protein
28	114	82.6	168	24 ABR39081	Human ovarian anti
29	114	82.6	201	23 ABA41630	bcl-x(L)/bcl-2 ass
30	113	81.9	23	17 AAR95166	Mutant Bcl2 compet
31	111	80.4	25	23 ABG78490	Human Bad peptide
32	111	80.4	25	23 ABG78490	Mutant Bcl2 compet
33	110	79.7	25	23 ABG78488	Mutant Bcl2 compet
34	110	79.7	25	23 ABG78489	Mutant Bcl2 compet
35	110	79.7	25	23 AAU78615	Human Bad peptide
36	110	79.7	25	23 AAU78616	Human Bad peptide
37	109	79.0	23	23 AAU78628	Human Bad peptide
38	109	79.0	25	23 ABG78486	Mutant Bcl2 compet
39	109	79.0	25	23 ABG78492	Mutant Bcl2 compet
40	109	79.0	25	23 ABG78497	Mutant Bcl2 compet
41	109	79.0	25	23 AAU78612	Human Bad peptide
42	109	79.0	25	23 AAU78619	Human Bad peptide
43	109	79.0	25	23 AAU78624	Human Bad peptide
44	108	78.3	25	23 ABG78485	Mutant Bcl2 compet
45	108	78.3	25	23 AAU78611	Human Bad peptide

ALIGNMENTS

RESULT 1

AAB37001
ID AAB37001 standard; peptide; 26 AA.
XX AAB37001;
AC AAB37001;
XX
XX 28-FEB-2001 (first entry)
DT Bcl2 polypeptide BH3 domain peptide #1.
DE
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.
XX
OS Homo sapiens.
XX
XX WC200059526-AL.
XX
XX 12-OCT-2000.
XX
XX 06-APR-2000; 2000WO-US09352.
XX
XX 07-APR-1999; 99US-0128202.
XX
XX (UJVE-) UNIV JEFFERSON THOMAS.
XX
XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX WPI; 2000-679325/66.
DR

Seq UPDATE 1-3, 55-56

(328-29 - Not Reported)

Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
WPI: 2000-679325/66.
New peptide conjugates for modulating apoptosis or for inhibiting B cell lymphoma/leukemia 2 (Bcl-2) function especially useful for treating neurodegenerative disorders, stroke, or cancer -
Claim 18: Page 17: 74pp: English.
The invention relates to a peptide conjugate having the formula:
(R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached to the N-terminus of the peptide, or a side chain of the peptide where the functional group of the side chain is NH₂ or OH; or X = O or NH, when the R-X group is attached to the C-terminus of the peptide, or a side chain of the peptide, where the side chain functional group is COOH or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C cyclohexyl containing one or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally monosubstituted with a 1-5C straight or branched chain alkyl group, phenyl optionally monosubstituted with a 1-5C straight or branched chain alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples of the peptide portion of the conjugate. The peptides represent analogues of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of the BH3 domain of the cell death agonist bcl-2. The peptide conjugate is useful for modulating apoptosis in the cells of a subject, or for reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of apoptosis in cancer cells. It is also useful for inhibiting Bcl-2 function. In particular, the peptide conjugate is useful for treating a subject afflicted with a cancer characterized by cancer cells that express Bcl-2. The cancer includes prostate, colorectal, gastric, non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide conjugate is also useful for treating disorders characterized by increased apoptosis, e.g. neurodegenerative disorders, acquired immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

Query Match	100.0%;	Score 138;	DB 21;	Length 26;
Best Local Similarity	100.0%;	Pred. No. 2e-14;		
Matches 26;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	NLWAAQRYGRELRRMSDFEGSFKGL	26	
Db	1	NLWAAQRYGRELRRMSDFEGSFKGL	26	
RESULT 3				
AAB37003				
ID	AAB37003	standard; peptide; 27 AA.		
XX	XX	AC	XX	
XX	AAB37003;			
DT	28-FEB-2001	(first entry)		
XX	XX			
DE	Bcl2 polypeptide BH3 domain peptide #3.			
XX	XX			
KW	Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;			
KW	cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;			
KW	apoptosis modulation; B cell lymphoma/Leukemia 2; cancer; prostate;			
KW	colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;			
KW	melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;			
XX	stroke; myocardial infarction.			
XX				
XX	Homo sapiens.			
XX	OS			
XX	XX			
FN	WO200059526-A1.			
XX	XX			
XX	XX			
PD	12-OCT-2000.			
XX	XX			
XX	06-APR-2000; 2000WO-US09352.			
PF	PF			
XX	XX			
XX	07-APR-1999; 99US-Q128202.			
PR	PR			

XX PA (UYJE-) UNIV JEFFERSON THOMAS.

XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX XX WPI; 2000-679325/66.

XX PT New peptide conjugates for modulating apoptosis or for inhibiting B

XX PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for

XX PT treating neurodegenerative disorders, stroke, or cancer

XX PS Claim 18; Page 17; 74pp; English.

XX CC The invention relates to a peptide conjugate having the formula:

CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached

CC to the N-terminus of the peptide, or a side chain of the peptide where

CC the functional group of the side chain is NH2 or OH; or X = O or NH,

CC when the R-X group is attached to the C-terminus of the peptide, or a

CC side chain of the peptide, where the side chain functional group is COOH

CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one

CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally

CC monosubstituted with a 1-5C straight or branched chain alkyl group,

CC phenyl optionally monosubstituted with a 1-5C straight or branched chain

CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples

CC of the peptide portion of the conjugate. The peptides represent analogues

CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of

CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is

CC useful for modulating apoptosis in the cells of a subject, or for

CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2

CC function. In particular, the peptide conjugate is useful for treating a

CC subject afflicted with a cancer characterized by cancer cells that

CC express Bcl-2. The cancer includes prostate, colorectal, gastric,

CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or

CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide

CC conjugate is also useful for treating disorders characterized by

CC increased apoptosis, e.g. neurodegenerative disorders, acquired

CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX CC Sequence 27 AA;

XX SQ Query Match 100.0%; Score 138; DB 21; Length 27;

XX Best Local Similarity 100.0%; Pred. No. 2.1e-14;

XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSFKGL 26

DB 1 NLWAAQRYGRELRRMSDEFGSFKGL 26

RESULT 4

AAB37056

ID AAB37056 standard; peptide; 27 AA.

XX AC AAB37056;

XX DT 28-FEB-2001 (first entry)

XX DE Bcl2 polypeptide BH3 domain peptide #56.

XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;

XX KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;

XX KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;

XX KW melanoma; lymphocytic leukemia; renal; thyroid; neuroblastoma;

XX KW stroke; myocardial infarction.

XX OS Homo sapiens.

XX XX WO200059526-A1.

XX PD 12-OCT-2000.

XX PN

PF 06-APR-2000; 2000WO-US09352.

XX PR 07-APR-1999; 99US-0128202.

XX PA (UYJE-) UNIV JEFFERSON THOMAS.

XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX XX WPI; 2000-679325/66.

XX PT New peptide conjugates for modulating apoptosis or for inhibiting B

XX PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for

XX PT treating neurodegenerative disorders, stroke, or cancer

XX PS Claim 18; Page 19; 74pp; English.

XX CC The invention relates to a peptide conjugate having the formula:

CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached

CC to the N-terminus of the peptide, or a side chain of the peptide where

CC the functional group of the side chain is NH2 or OH; or X = O or NH,

CC when the R-X group is attached to the C-terminus of the peptide, or a

CC side chain of the peptide, where the side chain functional group is COOH

CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one

CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally

CC monosubstituted with a 1-5C straight or branched chain alkyl group,

CC phenyl optionally monosubstituted with a 1-5C straight or branched chain

CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples

CC of the peptide portion of the conjugate. The peptides represent analogues

CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of

CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is

CC useful for modulating apoptosis in the cells of a subject, or for

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CC function. In particular, the peptide conjugate is useful for treating a

CC subject afflicted with a cancer characterized by cancer cells that

CC express Bcl-2. The cancer includes prostate, colorectal, gastric,

CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or

CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide

CC conjugate is also useful for treating disorders characterized by

CC increased apoptosis, e.g. neurodegenerative disorders, acquired

CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX CC Sequence 27 AA;

XX SQ Query Match 100.0%; Score 138; DB 21; Length 27;

XX Best Local Similarity 100.0%; Pred. No. 2.1e-14;

XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSFKGL 26

DB 2 NLWAAQRYGRELRRMSDEFGSFKGL 27

RESULT 5

AAB37055

ID AAB37055 standard; peptide; 28 AA.

XX AC AAB37055;

XX DT 28-FEB-2001 (first entry)

XX DE Bcl2 polypeptide BH3 domain peptide #55.

XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;

XX KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;

XX KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;

XX KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;

XX KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;

XX KW stroke; myocardial infarction.

XX OS Homo sapiens.

XX XX WO200059526-A1.

XX PN

XX 12-OCT-2000.
 PD
 XX 06-APR-2000; 2000WO-US09352.
 PF
 XX 07-APR-1999; 99US-0128202.
 PR
 XX (UKJE-) UNIV JEFFERSON THOMAS.
 PA
 XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 PI
 XX WPI; 2000-679325/66.
 DR
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 XX
 XX Claim 18; Page 19; 74pp; English.
 PS
 XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)n-peptide where n = 1-10; X = C-O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB7001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
 XX
 XX Sequence 28 AA;
 SQ
 Query Match 100.0%; Score 138; DB 21; Length 28;
 Best Local Similarity 100.0%; Pred. No. 2.1e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDEFGSFKGL 26
 Db 2 NLWAAQRYGRELRRMSDEFGSFKGL 27
 RESULT 6
 AAB70370
 ID AAB70370 standard; protein; 162 AA.
 XX
 XX AAB70370;
 AC
 XX
 XX 02-MAY-2001 (first entry)
 DT
 XX Shorter murine BAD mutant amino acid sequence SEQ ID NO:3.
 DE
 XX Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 KW immunostimulant; neuroprotective; neurotropic; antiischaemic; vulnary;
 KW cytostatic; antiviral; antiarthritic; antiinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.

XX Mus musculus.
 OS Synthetic.
 OS
 XX WO200110888-A1.
 PN
 XX 15-FEB-2001.
 PD
 XX 30-MAY-2000; 2000WO-US11864.
 PF
 XX 28-MAY-1999; 99US-0136783.
 PR
 XX (APOP-) APOPTOSIS TECHNOLOGY INC.
 PA
 XX Zhou X;
 PI
 XX WPI; 2001-138734/14.
 DR
 XX New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -
 PS
 XX Claim 7; Page 148-149; 157pp; English.
 XX The present invention describes an isolated or synthetic polypeptide
 CC (I) comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
 CC neurotropic, antiischaemic, vulnary, cytostatic, antiviral,
 CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed shorter murine BAD mutant amino acid sequence from the present
 CC invention.
 XX
 XX Sequence 162 AA;
 SQ
 Query Match 100.0%; Score 138; DB 22; Length 162;
 Best Local Similarity 100.0%; Pred. No. 1.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDEFGSFKGL 26
 Db 98 NLWAAQRYGRELRRMSDEFGSFKGL 123
 RESULT 7
 AAR95168
 ID AAR95168 standard; Protein; 204 AA.
 XX
 XX AAR95168;
 AC
 XX
 XX 06-JAN-1997 (first entry)
 DT
 XX bcl-x(L)/bcl-2 associated death promoter protein.
 DE
 XX Epitope; murine; bcl-x(L)/bcl-2 associated death promoter; Bad; stroke;
 KW polypeptide; bcl-x; cell death; regulate; BH1; BH2; apoptotic cell death;
 KW cytokine deprivation; IL-3 dependent cell line; immunodeficiency; AIDS;
 KW neurodegenerative disease; senescence; ischaemia; neoplasia.
 XX
 XX Mus musculus.
 OS

PH Key Location/Qualifiers
 FT Region 147..149
 FT /note="BH1 conserved amino acids"
 FT Region 191..192
 FT /note="BH2 conserved amino acids"
 FT Domain 38..61
 FT /note="PEST sequence"
 FT Domain 111..130
 FT /note="PEST sequence"
 XX WO9613614-A1.
 PN
 XX
 XX 09-MAY-1996.
 XX
 XX 31-OCT-1995; 95WO-US14246.
 XX
 XX 31-OCT-1994; 94US-0333565.
 XX
 XX (UNIW) UNIV WASHINGTON.
 XX
 XX Korsmeyer SJ;
 XX
 XX WPI: 1996-251465/25.
 XX N-PSDB; AAT29479.
 XX
 XX Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
 FT useful to treat neoplasia and apoptosis and to identify agents
 FT inhibiting its binding to bcl-2 or bcl-x(L) to form heteromultimers
 XX
 XX Claim 3; Fig 1; 130pp; English.
 XX
 XX This sequence represents the murine bcl-x(L)/bcl-2 associated death
 CC promoter (Bad) gene. Bad is a 22.1 kD protein which interacts with
 CC bcl-2 and bcl-x proteins and regulates cell death. It has homology
 CC to the bcl-2-related family clustered in the BH1 and BH2 domain. Bad
 CC has been found to hybridise to bcl-x(L) and bcl-2 in yeast two-hybrid
 CC assays and in vivo in mammalian cells. Overexpressed Bad counters the
 CC death inhibitory activity of bcl-x(L), but is much less effective at
 CC countering the death inhibitory activity of bcl-2. Bad expression can
 CC accelerate apoptotic cell death induced by cytokine deprivation in an
 CC IL-3 dependent cell line expressing bcl-x(L), and its also counters the
 CC death repressor activity of bcl-x(L). Bad competes with Bax for binding
 CC to bcl-x(L). Bad may be used to identify agents which inhibit its
 CC binding to bcl-2 or bcl-x(L) to form heterodimers. Such agents may be
 CC used to treat neurodegenerative diseases, immunodeficiency diseases,
 CC e.g. AIDS, senescence or ischaemia.
 XX
 XX Sequence 204 AA;
 SQ
 Query Match 100.0%; Score 138; DB 17; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.9e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
 |||||
 DB 140 NLWAAQRYGRELRRMSDFEGSFKGL 165
 RESULT 8
 AAW61315
 ID AAW61315 standard; Protein; 204 AA.
 XX
 XX AAW61315;
 XX
 XX 07-OCT-1998 (first entry)
 XX
 XX Murine BCL-XL/BCL-2 associated cell death regulator.
 DE
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 XX serine substituted mutant; apoptosis; cancer; viral infection.
 KW
 XX Mus sp.
 OS
 XX Mus sp.
 XX
 XX WO9613614-A1.
 PN

PM WO9817682-A1.
 XX
 XX 30-APR-1998.
 XX
 XX 17-OCT-1997; 97WO-US19175.
 XX
 XX 18-OCT-1996; 96US-0733505.
 XX
 XX (UNIW) UNIV WASHINGTON.
 XX
 XX Korsmeyer SJ;
 XX
 XX WPI: 1998-361422/23.
 XX N-PSDB; AAV27833.
 XX
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 FT useful for, e.g. treating reduced apoptosis such as in cancer or
 FT viral infection
 XX
 XX Claim 1; Fig 10; 95pp; English.
 XX
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence is the murine BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX
 XX Sequence 204 AA;
 SQ
 Query Match 100.0%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.9e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
 |||||
 DB 140 NLWAAQRYGRELRRMSDFEGSFKGL 165
 RESULT 9
 AAW61316
 ID AAW61316 standard; Protein; 204 AA.
 XX
 XX AAW61316;
 XX
 XX 07-OCT-1998 (first entry)
 XX
 XX Mutant BCL-XL/BCL-2 associated cell death regulator #1.
 DE
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 XX serine substituted mutant; apoptosis; cancer; viral infection.
 KW
 XX Mus sp.
 OS
 XX Synthetic.
 XX
 XX WO9817682-A1.
 PN

```

XX PD 30-APR-1998.
XX PF 17-OCT-1997; 97WO-US19175.
XX PF 18-OCT-1996; 96US-0733505.
XX PR 18-OCT-1996; 96US-0733505.
XX PA (UNIW ) UNIV WASHINGTON.
XX PA Korsmeyer SJ;
XX PI WPI: 1998-261422/23.
XX DR N-PSDB; AAV27834.
XX DR New mutant BAD polypeptide with phosphorylatable serine replaced -
XX PT useful for, e.g. treating reduced apoptosis such as in cancer or
XX PT viral infection
XX PS Claim 7; Page 59; 95pp; English.
XX CC The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
XX CC death regulator) proteins, having an amino acid other than Ser at
XX CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
XX CC present sequence represents a mutant BAD protein. Also described are: (1)
XX CC fragments of mutant BAD protein able to decrease cell viability; (2)
XX CC fusion proteins of mutant BAD with a heterologous polypeptide that
XX CC increases intracellular delivery. Mutant BAD proteins are used to treat
XX CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
XX CC viral infection, lymphoproliferation, arthritis, infertility,
XX CC inflammation and autoimmune disease. Polynucleotide sequences encoding
XX CC mutant BAD proteins can be used similarly by gene therapy or to produce
XX CC transgenic animals for use as disease models or in drug screening. BAD
XX CC proteins phosphorylated at specified Ser are used to screen for enhancers
XX CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
XX CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
XX CC aging or ischaemic cell death. The apoptotic status of cells is
XX CC determined by measuring relative amounts of phosphorylated and non-
XX CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
XX CC greater death-promoting activity than wild-type BAD which can become
XX CC phosphorylated on the specified Ser, forming a product that does not
XX CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
XX CC proteins in the cytosol, thus promoting cell survival. The mutants with
XX CC Ser substituted cannot bind 14-3-3.
XX SQ Sequence 204 AA;
XX Query Match 100.0%; Score 138; DB 19; Length 204;
XX Best Local Similarity 100.0%; Pred. No. 1.9e-13;
XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 NLWAAQRYGRELRLMSDEFGSFGL 26
DB 140 NLWAAQRYGRELRLMSDEFGSFGL 165
RESULT 10
AAW61317
ID AAW61317 standard; Protein; 204 AA.
XX AC AAW61317;
XX DT 07-OCT-1998 (first entry)
XX DE Mutant BCL-XL/BCL-2 associated cell death regulator #2.
XX KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
XX KW serine substituted mutant; apoptosis; cancer; viral infection.
XX OS Mus sp.
XX OS Synthetic.
XX PN WO9817682-A1.
XX PD 30-APR-1998.

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PD 30-APR-1998.
XX 17-OCT-1997; 97WO-US19175.
XX 18-OCT-1996; 96US-0733505.
XX (UNIW ) UNIV WASHINGTON.
XX Korsmeyer SJ;
XX WPI: 1998-261422/23.
XX N-PSDB; AAV27835.
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
XX useful for, e.g. treating reduced apoptosis such as in cancer or
XX viral infection
XX Claim 7; Page 60; 95pp; English.
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
XX death regulator) proteins, having an amino acid other than Ser at
XX position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
XX present sequence represents a mutant BAD protein. Also described are: (1)
XX fragments of mutant BAD protein able to decrease cell viability; (2)
XX fusion proteins of mutant BAD with a heterologous polypeptide that
XX increases intracellular delivery. Mutant BAD proteins are used to treat
XX or prevent diseases associated with reduced apoptosis, e.g. cancer,
XX viral infection, lymphoproliferation, arthritis, infertility,
XX inflammation and autoimmune disease. Polynucleotide sequences encoding
XX mutant BAD proteins can be used similarly by gene therapy or to produce
XX transgenic animals for use as disease models or in drug screening. BAD
XX proteins phosphorylated at specified Ser are used to screen for enhancers
XX and inhibitors of serine-phosphatase. Inhibitors are potentially useful
XX in treatment of excessive apoptosis such as AIDS, neurodegeneration,
XX aging or ischaemic cell death. The apoptotic status of cells is
XX determined by measuring relative amounts of phosphorylated and non-
XX phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
XX greater death-promoting activity than wild-type BAD which can become
XX phosphorylated on the specified Ser, forming a product that does not
XX heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
XX proteins in the cytosol, thus promoting cell survival. The mutants with
XX Ser substituted cannot bind 14-3-3.
XX SQ Sequence 204 AA;
XX Query Match 100.0%; Score 138; DB 19; Length 204;
XX Best Local Similarity 100.0%; Pred. No. 1.9e-13;
XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 NLWAAQRYGRELRLMSDEFGSFGL 26
DB 140 NLWAAQRYGRELRLMSDEFGSFGL 165
RESULT 11
AAW61318
ID AAW61318 standard; Protein; 204 AA.
XX AC AAW61318;
XX DT 07-OCT-1998 (first entry)
XX DE Mutant BCL-XL/BCL-2 associated cell death regulator #3.
XX KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
XX KW serine substituted mutant; apoptosis; cancer; viral infection.
XX OS Mus sp.
XX OS Synthetic.
XX PN WO9817682-A1.
XX PD 30-APR-1998.

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XX PF 17-OCT-1997; 97WO-US19175.
XX PR 18-OCT-1996; 96US-0733505.
XX PA (UNTW ) UNIV WASHINGTON.
XX PI Kormsmeier SJ;
XX WPI: 1998-261422/23.
XX N-PSDB; RAV27836.
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
XX PT useful for, e.g. treating reduced apoptosis such as in cancer or
XX PT viral infection.
XX
XX Claim 7; Page 60-61; 95pp; English.
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
XX death regulator) proteins, having an amino acid other than Ser at
XX position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
XX present sequence represents a mutant BAD protein. Also described are: (1)
XX fragments of mutant BAD protein able to decrease cell viability; (2)
XX fusion proteins of mutant BAD with a heterologous polypeptide that
XX increases intracellular delivery. Mutant BAD proteins are used to treat
XX or prevent diseases associated with reduced apoptosis, e.g. cancer,
XX viral infection, lymphoproliferation, arthritis, infertility,
XX inflammation and autoimmune disease. Polynucleotide sequences encoding
XX mutant BAD proteins can be used similarly by gene therapy or to produce
XX transgenic animals for use as disease models or in drug screening. BAD
XX proteins phosphorylated at specified Ser are used to screen for enhancers
XX and inhibitors of serine-phosphatase. Inhibitors are potentially useful
XX in treatment of excessive apoptosis such as AIDS, neurodegeneration,
XX aging or ischaemic cell death. The apoptotic status of cells is
XX determined by measuring relative amounts of phosphorylated and non-
XX phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
XX greater death-promoting activity than wild-type BAD which can become
XX phosphorylated on the specified Ser, forming a product that does not
XX heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
XX proteins in the cytosol, thus promoting cell survival. The mutants with
XX Ser substituted cannot bind 14-3-3.
XX
XX SQ Sequence 204 AA;
    Query Match 100.0%; Score 138; DB 19; Length 204;
    Best Local Similarity 100.0%; Pred. No. 1.9e-13;
    Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
   |||||
Db 140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 12
AAW58832
ID AAW58832 standard; protein; 204 AA.
XX
XX AAW58832;
XX
XX 23-JUL-1998 (first entry)
XX
XX Murine BAD protein.
XX
XX BAD protein; Bcl-XL/Bcl-2 associated cell death regulator; 14-3-3;
XX serine phosphorylation; post-translational modification; apoptosis;
XX signal transduction regulator; phosphoserine phosphatase; senescence;
XX immunodeficiency disease; neurodegenerative disease; infertility;
XX cancer, viral infection; lymphoproliferative condition; arthritis;
XX inflammation; autoimmune diseases.
XX
XX Mus sp.
XX
XX WO9809643-A1.

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XX 12-MAR-1998.
XX
XX 09-SEP-1997; 97WO-US15871.
XX
XX 09-SEP-1996; 96US-0707868.
XX
XX (UNIW ) UNIV WASHINGTON.
XX
XX Kormsmeier SJ;
XX
XX WPI: 1998-207049/18.
XX
XX Serine-phosphorylated Bcl-XL/Bcl-2 Associated cell Death regulator
XX polypeptide - useful for modulation of apoptosis associated with,
XX e.g. cancer and immunodeficiency diseases
XX
XX Claim 3; Fig 8; 61pp; English.
XX
XX This sequence represents a novel serine-phosphorylated protein, BAD
XX (Bcl-XL/Bcl-2 associated cell death regulator). The serine residue is
XX phosphorylated in a post-translational modification and allows binding
XX to the 14-3-3 protein which is a signal transduction regulator.
XX Modulators of phosphorylated BAD, which act through inhibition/activation
XX of a phosphoserine phosphatase, are useful for preventing/treating
XX increased/decreased apoptosis in a cell. The increased apoptosis may
XX result from immunodeficiency diseases, senescence, neurodegenerative
XX disease, ischaemic cell death, reperfusion cell death, infertility and
XX wound-healing. Decreased apoptosis may result from cancer, viral
XX infection, lymphoproliferative conditions, arthritis, infertility,
XX inflammation and autoimmune diseases. Measuring the amount of
XX phosphorylated compared to unphosphorylated BAD polypeptide and/or total
XX BAD in a cell is useful for determining the apoptotic state of a cell.
XX
XX SQ Sequence 204 AA;
    Query Match 100.0%; Score 138; DB 19; Length 204;
    Best Local Similarity 100.0%; Pred. No. 1.9e-13;
    Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
   |||||
Db 140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 13
AAW570369
ID AAW570369 standard; protein; 204 AA.
XX
XX AAW570369;
XX
XX 02-MAY-2001 (first entry)
XX
XX Longer murine BAD mutant amino acid sequence SEQ ID NO:2.
XX
XX Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
XX immunostimulant; neuroprotective; nootropic; antischismatic; vulnary;
XX cytostatic; antiviral; antiarthritic; antiinflammatory; wound healing;
XX immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
XX immunodeficiency disease; neurodegenerative disease; viral infection;
XX ischaemic cell death; reperfusion cell death; arthritis; infertility;
XX lymphoproliferative condition; inflammation; autoimmune disease.
XX
XX Mus musculus.
XX
XX Synthetic.
XX
XX WO200110888-A1.
XX
XX 15-FEB-2001.
XX
XX 30-MAY-2000; 2000WO-US11864.
XX
XX 28-MAY-1999; 99US-0136783.

```

XX PA (APOP-) APOPTOSIS TECHNOLOGY INC.
 XX PT Zhou X;
 XX XX WPI; 2001-138734/14.
 XX DR
 XX PT New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 XX PT useful for screening for candidate compounds which induce or inhibit
 XX PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 XX PT Ser113.
 XX XX
 XX PS Claim 4; Page 148; 157pp; English.
 XX CC
 XX CC The present invention describes an isolated or synthetic polypeptide
 XX CC (1) comprising a less than full length amino acid sequence of a mutant
 XX CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 XX CC fragment, which contains amino acid substitutions at Ser118 of a human
 XX CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 XX CC BAD (shorter murine BAD). (1) has immunostimulant, neuroprotective,
 XX CC neurotropic, antiischaemic, vulnerary, cytotstatic, antiviral,
 XX CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 XX CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 XX CC polynucleotides can be used for screening candidate compounds and drugs
 XX CC for activity that promote cell survival or apoptosis. Other uses include
 XX CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 XX CC identified and (mutant) BAD polypeptides are useful in treating
 XX CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 XX CC death, reperfusion cell death, wound healing, cancer, viral infections,
 XX CC lymphoproliferative conditions, arthritis, infertility, inflammation, and
 XX CC autoimmune diseases. The present sequence represents a specifically
 XX CC claimed longer murine BAD mutant amino acid sequence from the present
 XX CC invention.
 XX SQ
 XX SQ Sequence 204 AA;
 Query Match 100.0%; Score 138; DB 22; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.9e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRLMSDEFGSGKGL 26
 Db 140 NLWAAQRYGRELRLMSDEFGSGKGL 165
 RESULT 14
 ABR39082
 ID ABR39082 standard; Protein; 204 AA.
 XX ABR39082;
 XX AC
 XX AC
 XX DT 10-MAY-2003 (first entry)
 XX DE
 XX DE Murine BAD protein SEQ ID NO:4.
 XX KW Murine; BAD; herpes simplex virus; HSV; US3; herpes virus; apoptosis;
 XX KW virucide; infection.
 XX OS
 XX OS Mus musculus.
 XX PN
 XX PN WO2003012049-A2.
 XX XX
 XX PD 13-FEB-2003.
 XX PF
 XX PF 31-JUL-2002; 2002WO-US24177.
 XX PR
 XX PR 31-JUL-2001; 2001US-308999P.
 XX XX
 XX XX (UYCH-) UNIV CHICAGO.
 XX XX
 XX PI Munger J, Roizman B;
 XX XX
 XX XX WPI; 2003-248168/24.
 XX DR

DR N-PSDB; AB281201.
 XX
 XX PT Inducing apoptosis in a cell infected with herpes simplex virus, HSV,
 XX PT by administering to the cell, a composition comprising an agent that
 XX PT inhibits phosphorylation of pro-apoptotic polypeptide BAD by HSV US3.
 XX XX
 XX PS Claim 15; Page 168; 192pp; English.
 XX CC
 XX CC The present invention describes a method (M1) for inducing apoptosis in
 XX CC a cell infected with herpes simplex virus (HSV), which comprises:
 XX CC administering to the cell, a composition having an agent that inhibits
 XX CC phosphorylation of pro-apoptotic polypeptide BAD by HSV US3. Also
 XX CC described is a method (M2) for treating a patient infected with HSV, by
 XX CC administering to the patient, a composition comprising a peptide
 XX CC comprising a sequence of 4-100 continuous amino acids of a 168 residue
 XX CC amino acid sequence (see ABR39081), where the peptide comprises ser112,
 XX CC ser135 or ser155, or their combinations. BAD has virucide activity.
 XX CC M1 is useful for inducing apoptosis in a cell infected with HSV, where
 XX CC the cell is in a human. M2 is useful for treating a patient infected
 XX CC with HSV. The present sequence represents murine BAD, which is used in
 XX CC the exemplification of the present invention.
 XX SQ
 XX SQ Sequence 204 AA;
 Query Match 100.0%; Score 138; DB 24; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.9e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRLMSDEFGSGKGL 26
 Db 140 NLWAAQRYGRELRLMSDEFGSGKGL 165
 RESULT 15
 AAU00220
 ID AAU00220 standard; Protein; 567 AA.
 XX AAU00220;
 XX AC
 XX AC
 XX DT 31-MAY-2001 (first entry)
 XX DE
 XX DE Bad-DTRR apoptosis-modifying fusion protein.
 XX KW Mouse; Bad-DTRR; apoptosis; cancer; spinal muscular atrophy;
 XX KW diphtheria toxin receptor binding domain; DTR; neoplasia; tumour;
 XX KW hyper-proliferation; Alzheimer's disease; neurodegenerative disorder;
 XX KW transient ischaemic neuronal injury; stroke; spinal cord injury;
 XX KW Huntington's disease.
 XX OS
 XX OS Chimeric - Mus sp.
 XX OS Chimeric - Corynebacterium diphtheriae.
 XX OS Chimeric - Synthetic.
 XX FH
 XX FH Key Location/Qualifiers
 XX FT Region 3..12
 XX FT /note= "10x histidine tag"
 XX XX
 XX PN WO200112661-A2.
 XX XX
 XX XX 22-FEB-2001.
 XX XX
 XX XX 15-AUG-2000; 2000WO-US22293.
 XX XX
 XX XX 16-AUG-1999; 99US-0149220.
 XX XX
 XX XX (HARD) HARVARD COLLEGE.
 XX XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX XX
 XX XX Youle RJ, Liu X, Collier RJ;
 XX XX
 XX XX WPI; 2001-218343/22.
 XX XX N-PSDB; AAS00248.
 XX DR
 XX DR

PT Novel fusion protein for modifying apoptosis in target cell and
 PT reducing apoptosis after transient ischaemic neuronal injury, has two
 PT domains which targets protein to a cell and modifies apoptotic response
 PT of cell -

XX
 XX
 PS Claim 4; Page 59-61; 65pp; English.

CC The sequence represents the amino acid sequence of Bad-DTRR apoptosis-
 CC modifying fusion protein comprising Bad gene sequence fused via a short
 CC linker to diphtheria toxin translocation domain (DTRR). The
 CC functional apoptosis-modifying fusion protein is capable of binding a
 CC target cell and integrating into or crossing a cellular membrane of the
 CC target cell. The apoptosis-modifying fusion protein comprises at least
 CC two domains: the DTR domain, which targets the fusion protein to the
 CC target cell and the Bcl-XL domain, which modifies an apoptotic response
 CC of the target cell. The fusion protein is useful for modifying
 CC (inhibiting or enhancing) apoptosis in a target cell, such as neuron,
 CC lymphocyte, cancer, neoplasm, macrophage, epithelial, stem, tumour or
 CC hyper-proliferative cell or an adipocyte. It is also useful for reducing
 CC apoptosis in a subject after transient ischaemic neuronal injury,
 CC especially spinal cord injury. The fusion protein may be used to treat
 CC various diseases and injury conditions through inhibition or enhancement
 CC of apoptotic cellular response, including neurodegenerative disorders
 CC such as Alzheimer's disease, Huntington's disease, spinal muscular
 CC atrophy, stroke episodes and unregulated cell growth as in tumours and
 CC various cancers. The apoptosis-modifying fusion protein can be delivered
 CC effectively throughout the body and targeted to selective tissue and
 CC cells.

XX
 SQ Sequence 567 AA;

Query Match 100.0%; Score 138; DB 22; Length 567;
 Best Local Similarity 100.0%; Pred. No. 6.1e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAQRYGRELRLMSDEFGSPKGL 26
 |||||
 Db 161 NLWAQRYGRELRLMSDEFGSPKGL 186

Search completed: September 15, 2003, 17:22:13
 Job time : 37.7714 secs

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:22:21 ; Search time 13.5571 Seconds
(without alignments)
81.144 Million cell updates/sec

Title: US-09-544-664-1

Perfect score: 138

Sequence: 1 NLWAAQRYGRELRRMSDFEGSPKGL 26

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	138	100.0	204	1	US-08-333-565-2
2	138	100.0	204	2	US-08-661-479-2
3	138	100.0	204	2	US-08-733-505A-1
4	138	100.0	204	2	US-08-733-505A-12
5	138	100.0	204	2	US-08-733-505A-13
6	138	100.0	204	2	US-08-733-505A-14
7	135	97.8	204	2	US-08-717-123-3
8	135	97.8	204	4	US-09-375-257-3
9	114	82.6	166	1	US-08-665-617-2
10	114	82.6	166	2	US-08-717-123-2
11	114	82.6	168	3	US-08-985-335-1
12	114	82.6	168	3	US-08-985-335-7
13	114	82.6	168	3	US-09-410-372-1
14	114	82.6	168	3	US-09-410-372-7
15	114	82.6	168	4	US-09-375-257-2
16	113	81.9	23	1	US-08-333-565-10
17	113	81.9	23	2	US-08-661-479-10
18	102	73.9	59	2	US-08-733-505A-55
19	102	73.9	59	2	US-08-733-505A-56
20	102	73.9	59	2	US-08-733-505A-57
21	102	73.9	59	1	US-08-733-505A-58
22	86	62.3	16	1	US-08-333-565-26
23	86	62.3	16	2	US-08-661-479-26
24	61	44.2	11	2	US-08-733-505A-34
25	61	44.2	11	2	US-08-706-741B-69
26	61	44.2	11	2	US-08-924-695A-69
27	51	37.0	66	2	US-08-667-087B-40

28 48.5 35.1 904 4 US-09-328-352-4656 Sequence 4656, Ap
29 46 33.3 610 4 US-09-252-991A-19594 Sequence 19594, A
30 46 33.3 946 3 US-09-074-573-3 Sequence 3, Appli
31 46 33.3 946 3 US-09-388-774-3 Sequence 3, Appli
32 46 33.3 946 4 US-09-546-153-1 Sequence 1, Appli
33 45.5 33.0 906 4 US-09-252-991A-31458 Sequence 31458, A
34 45 32.6 229 4 US-09-328-352-5164 Sequence 23807, A
35 45 32.6 303 4 US-09-328-352-5164 Sequence 5164, Ap
36 45 32.6 356 4 US-09-235-103-2 Sequence 2, Appli
37 45 32.6 356 4 US-09-235-103-4 Sequence 4, Appli
38 45 32.6 1064 4 US-09-252-991A-17508 Sequence 17508, A
39 44.5 32.2 903 4 US-09-252-991A-28775 Sequence 28775, A
40 44 31.9 125 4 US-09-328-352-7449 Sequence 7449, Ap
41 44 31.9 263 4 US-09-651-656-27 Sequence 27, Appl
42 44 31.9 263 4 US-09-650-855-27 Sequence 20, Appl
43 44 31.9 877 4 US-09-206-551-20 Sequence 27, Appl
44 44 31.9 1125 4 US-09-252-991A-18729 Sequence 18729, A
45 43.5 31.5 467 4 US-09-252-991A-18296 Sequence 18296, A

ALIGNMENTS

RESULT 1

US-08-333-565-2
; Sequence 2, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bel-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "Deduced amino acid sequence
; OTHER INFORMATION: of mouse BAD."
US-08-333-565-2

Query Match 100.0%; Score 138; DB 1; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.8e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGL 26
 |||||
 Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 2

US-08-661-479-2
 ; Sequence 2, Application US/08661479
 ; Patent No. 5834209
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, Stanley J.
 ; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
 ; TITLE OF INVENTION: REGULATOR
 ; NUMBER OF SEQUENCES: 59
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Townsend and Townsend Kourie and Crew
 ; STREET: 379 Lytton Avenue
 ; CITY: Palo Alto
 ; STATE: California
 ; COUNTRY: US
 ; ZIP: 94301
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/661,479
 ; FILING DATE: 11-JUN-1995
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 08/333,565
 ; FILING DATE: 31-OCT-1994
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Smith, William M
 ; REGISTRATION NUMBER: 30,223
 ; REFERENCE/DOCKET NUMBER: 15726A-000700
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (415) 326-2400
 ; TELEFAX: (415) 326-2422
 ; INFORMATION FOR SEQ ID NO: 2:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 204 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: Protein
 ; FEATURE:
 ; NAME/KEY: Protein
 ; LOCATION: 1..204
 ; OTHER INFORMATION: /note= "Deduced amino acid sequence
 ; OF mouse BAD."
 ; US-08-661-479-2

Query Match 100.0%; Score 138; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 5.8e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGL 26
 |||||
 Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 3

US-08-733-505A-1
 ; Sequence 1, Application US/08733505A
 ; Patent No. 5856445
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, STANLEY J.
 ; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 ; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 ; NUMBER OF SEQUENCES: 60
 ; CORRESPONDENCE ADDRESS:

ADDRESSEE: HOWELL & HAFFERKAMP, L.C.
 STREET: 7733 FORSYTH BLVD., SUITE 1400
 CITY: ST. LOUIS
 STATE: MISSOURI
 COUNTRY: USA
 ZIP: 63105
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/733,505A
 ; FILING DATE:

CLASSIFICATION: 530
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: HOLLAND, DONALD R.
 ; REGISTRATION NUMBER: 35,197
 ; REFERENCE/DOCKET NUMBER: 965458
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314) 727-5188
 ; TELEFAX: (314) 727-6092
 ; INFORMATION FOR SEQ ID NO: 1:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 204 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS:
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-733-505A-1

Query Match 100.0%; Score 138; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 5.8e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGL 26
 |||||
 Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 4

US-08-733-505A-12
 ; Sequence 12, Application US/08733505A
 ; Patent No. 5856445
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, STANLEY J.
 ; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 ; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 ; NUMBER OF SEQUENCES: 60
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: HOWELL & HAFFERKAMP, L.C.
 ; STREET: 7733 FORSYTH BLVD., SUITE 1400
 ; CITY: ST. LOUIS
 ; STATE: MISSOURI
 ; COUNTRY: USA
 ; ZIP: 63105
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/733,505A
 ; FILING DATE:
 ; CLASSIFICATION: 530
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: HOLLAND, DONALD R.
 ; REGISTRATION NUMBER: 35,197
 ; REFERENCE/DOCKET NUMBER: 965458
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314) 727-5188
 ; TELEFAX: (314) 727-6092
 ; INFORMATION FOR SEQ ID NO: 12:

```

; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-12

Query Match          100.0%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.8e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
Db 140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 5
US-08-733-505A-13
; Sequence 13, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-13

Query Match          100.0%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.8e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
Db 140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 6
US-08-733-505A-14
; Sequence 14, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.

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; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-14

Query Match          100.0%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.8e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
Db 140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 7
US-08-717-123-3
; Sequence 3, Application US/08717123
; Patent No. 5965703
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
; Acids and Methods of Use
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE: 20-SEP-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815

```



```
/ REFERENCE/DOCKET NUMBER: P-ID 1929
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (619) 535-9001
/ TELEFAX: (619) 535-8949
/ INFORMATION FOR SEQ ID NO: 3:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 204 amino acids
/ TYPE: amino acid
/ TOPOLOGY: linear
US-08-717-123-3

Query Match          97.8%; Score 135; DB 2; Length 204;
Best Local Similarity 96.2%; Pred. No. 1.7e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGL 26
DB 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 8
US-09-375-257-3
; Sequence 3, Application US/09375257
; Patent No. 6504022
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D1
; CURRENT APPLICATION NUMBER: US/09/375,257
; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-375-257-3

Query Match          97.8%; Score 135; DB 4; Length 204;
Best Local Similarity 96.2%; Pred. No. 1.7e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGL 26
DB 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 9
US-08-665-617-2
; Sequence 2, Application US/08665617
; Patent No. 5663316
; GENERAL INFORMATION:
; APPLICANT: Xudong, Yin
; TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Saliwanchik & Saliwanchik
; STREET: 2421 N.W. 41st Street, Suite A-1
; CITY: Gainesville
; STATE: Florida
; COUNTRY: USA
; ZIP: 32606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/665,617
; FILING DATE:
; CLASSIFICATION: 530
```

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/ ATTORNEY/AGENT INFORMATION:
/ NAME: Saliwanchik, David R.
/ REGISTRATION NUMBER: 31,794
/ REFERENCE/DOCKET NUMBER: CL-8
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (352) 375-8100
/ TELEFAX: (352) 372-5800
/ INFORMATION FOR SEQ ID NO: 2:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 166 amino acids
/ TYPE: amino acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/ MOLECULE TYPE: protein
US-08-665-617-2

Query Match          82.6%; Score 114; DB 1; Length 166;
Best Local Similarity 91.7%; Pred. No. 2.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSK 24
DB 101 NLWAAQRYGRELRRMSDEFDSEK 124

RESULT 10
US-08-717-123-2
; Sequence 2, Application US/08717123
; Patent No. 5965703
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/717,123
; FILING DATE: 20-SEP-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-ID 1929
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-717-123-2

Query Match          82.6%; Score 114; DB 2; Length 168;
Best Local Similarity 91.7%; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSK 24
DB 103 NLWAAQRYGRELRRMSDEFDSEK 126
```

```

RESULT 11
US-08-985-335-1
; Sequence 1, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; ADDRESS: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-845-4166
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
US-08-985-335-1

Query Match      82.6%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRELRRMSDEFEGSK 24
   |||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 12
US-08-985-335-7
; Sequence 7, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9

```

```

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-845-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
US-08-985-335-7

Query Match      82.6%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRELRRMSDEFEGSK 24
   |||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 13
US-09-410-372-1
; Sequence 1, Application US/09410372
; Patent No. 6281334
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:

```

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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
US-09-410-372-1
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```
Query Match 82.68; Score 114; DB 3; Length 168;
Best Local Similarity 91.78; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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```
QY 1 NLWAAQRYGRELRRMSDEFEFGSK 24
| | | | | | | | | | | | | | | |
Db 103 NLWAAQRYGRELRRMSDEFEVDSFK 126
```

RESULT 14

```
US-09-410-372-7
; Sequence 7, Application US/09410372
; Patent No. 6281334
```

```
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSER: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
```

```
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:
```

```
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
```

```
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
```

```
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
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```
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
US-09-410-372-7
```

```
Query Match 82.68; Score 114; DB 3; Length 168;
Best Local Similarity 91.78; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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```
QY 1 NLWAAQRYGRELRRMSDEFEFGSK 24
| | | | | | | | | | | | | | | |
Db 103 NLWAAQRYGRELRRMSDEFEVDSFK 126
```

RESULT 15

```
US-09-375-257-2
; Sequence 2, Application US/09375257
; Patent No. 6504022
```

```
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D1
; CURRENT APPLICATION NUMBER: US/09/375,257
; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-375-257-2
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Query Match 82.68; Score 114; DB 4; Length 168;
Best Local Similarity 91.78; Pred. No. 2.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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```
QY 1 NLWAAQRYGRELRRMSDEFEFGSK 24
| | | | | | | | | | | | | | | |
Db 103 NLWAAQRYGRELRRMSDEFEVDSFK 126
```

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Search completed: September 15, 2003, 17:45:05
Job time : 14.5571 secs
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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:25:56 ; Search time 20.6143 Seconds
(without alignments)
184.034 Million cell updates/sec

Title: US-09-544-664-1

Perfect score: 138
Sequence: 1 NLWAAQRYGRELRRMSDFEGSFKGL 26

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 541936 seqs, 145912426 residues

Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA:*

- 1: /cgn2.6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
- 2: /cgn2.6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
- 3: /cgn2.6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
- 4: /cgn2.6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
- 5: /cgn2.6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
- 6: /cgn2.6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
- 7: /cgn2.6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
- 8: /cgn2.6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
- 9: /cgn2.6/ptodata/1/pubpaa/US09_PUBCOMB.pep.*
- 10: /cgn2.6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
- 11: /cgn2.6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
- 12: /cgn2.6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
- 13: /cgn2.6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
- 14: /cgn2.6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
- 15: /cgn2.6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
- 16: /cgn2.6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
- 17: /cgn2.6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
- 18: /cgn2.6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	135	97.8	204	9 US-09-922-378-3	Sequence 3, Appli
2	135	97.8	204	14 US-10-066-179-3	Sequence 3, Appli
3	114	82.6	25	15 US-10-059-261-258	Sequence 258, App
4	114	82.6	168	9 US-09-922-378-2	Sequence 2, Appli
5	114	82.6	168	9 US-09-894-657-1	Sequence 1, Appli
6	114	82.6	168	9 US-09-894-657-7	Sequence 7, Appli
7	114	82.6	168	14 US-10-066-179-2	Sequence 2, Appli
8	71	51.4	15	15 US-10-174-105A-147	Sequence 147, App
9	50	36.2	215	15 US-10-156-761-9145	Sequence 9145, Ap
10	47	34.1	35	15 US-10-092-750-1	Sequence 1, Appli
11	47	34.1	138	15 US-10-092-750-241	Sequence 241, App
12	46	33.3	682	12 US-10-238-075-1077	Sequence 1077, Ap
13	46	33.3	946	9 US-09-828-423-3	Sequence 3, Appli
14	44	31.9	272	15 US-10-156-761-11541	Sequence 11541, A
15	44	31.9	426	9 US-09-815-242-5704	Sequence 5704, Ap

16	44	31.9	699	14	US-10-008-355-8	Sequence 8, Appli
17	44	31.9	705	9	US-09-815-242-12463	Sequence 12463, A
18	44	31.9	712	14	US-10-008-355-2	Sequence 2, Appli
19	44	31.9	877	12	US-10-369-294-20	Sequence 20, Appli
20	43	31.2	213	9	US-09-843-846-18	Sequence 18, Appli
21	43	31.2	232	10	US-09-881-752A-238	Sequence 238, App
22	43	31.2	270	11	US-09-934-455-162	Sequence 162, App
23	43	31.2	380	9	US-09-149-045-2	Sequence 2, Appli
24	43	31.2	380	15	US-10-166-359-2	Sequence 2, Appli
25	43	31.2	380	15	US-10-166-1113-2	Sequence 2, Appli
26	43	31.2	380	15	US-10-166-357-2	Sequence 2, Appli
27	43	31.2	380	15	US-10-166-372-2	Sequence 2, Appli
28	43	31.2	380	15	US-10-184-722-3	Sequence 3, Appli
29	43	31.2	380	15	US-10-251-385-62	Sequence 62, Appli
30	43	31.2	380	15	US-10-251-385-198	Sequence 198, App
31	43	31.2	380	15	US-10-225-567A-233	Sequence 233, App
32	43	31.2	543	15	US-10-156-761-13485	Sequence 13485, A
33	43	31.2	571	9	US-09-815-242-11813	Sequence 11813, A
34	43	31.2	582	10	US-09-331-631A-22	Sequence 22, Appli
35	43	31.2	640	9	US-09-989-722-501	Sequence 501, App
36	43	31.2	640	9	US-09-989-723-501	Sequence 501, App
37	43	31.2	640	9	US-09-989-279-501	Sequence 501, App
38	43	31.2	640	9	US-09-989-727-501	Sequence 501, App
39	43	31.2	640	10	US-09-989-731-501	Sequence 501, App
40	43	31.2	640	10	US-09-989-732-501	Sequence 501, App
41	43	31.2	640	10	US-09-991-073-501	Sequence 501, App
42	43	31.2	640	10	US-09-909-320-292	Sequence 292, App
43	43	31.2	640	10	US-09-990-442-501	Sequence 501, App
44	43	31.2	640	10	US-09-991-163-501	Sequence 501, App
45	43	31.2	640	10	US-09-993-604-501	Sequence 501, App

ALIGNMENTS

RESULT 1
US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 97.8%; Score 135; DB 9; Length 204;
Best Local Similarity 96.2%; Pred. No. 3.3e-12;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
|||||
Db 140 NLWAAQRYGRELRRMTDFEGSFKGL 165

RESULT 2
US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE

```

; ORGANISM: Homo sapiens
US-09-922-378--2

Query Match      82.6%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 3.4e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFESEK 24
Db      103 NLWAAQRYGRELRLMSDEFEVSFK 126
        ||| ||||| ||||| ||||| |||

RESULT 5
US-09-894-657-1
; Sequence 1, Application US/09894657
; Patent No. US3002098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <UNKNOWN>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-894-657-1

Query Match      82.6%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 3.4e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFESEK 24
Db      103 NLWAAQRYGRELRLMSDEFEVSFK 126
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RESULT 6
US-09-894-657-7

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; Sequence 7, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 7:
US-09-894-657-7

Query Match      82.6%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 3.4e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFGSKF 24
      |||
Db      103 NLWAAQRYGRELRLMSDEFGSKF 126

RESULT 7
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168

Query Match      82.6%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 3.4e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFGSKF 24
      |||
Db      103 NLWAAQRYGRELRLMSDEFGSKF 126

RESULT 7
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168

Query Match      82.6%; Score 114; DB 14; Length 168;
Best Local Similarity 91.7%; Pred. No. 3.4e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFGSKF 24
      |||
Db      103 NLWAAQRYGRELRLMSDEFGSKF 126

RESULT 8
US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US2003006852A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECIFIC
; FILE OF INVENTION: CONTEXT-INDEPENDENT ANTIBODIES COUPLED WITH DATABASE SEARCHING
; FILE REFERENCE: CST-138 CIF3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147

Query Match      51.4%; Score 71; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.00061;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      9 GRELRLMSDEFGS 22
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Db      1 GRELRLMSDEFGS 14

RESULT 9
US-10-156-761-9145
; Sequence 9145, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
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; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 9145
; LENGTH: 215
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-9145

Query Match      36.2%; Score 50; DB 15; Length 215;
Best Local Similarity 56.2%; Pred. No. 12;
Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY      6 QYGRGLRRMSDEFG 21
DB      108 ERWGGDLRRMRDEADG 123

RESULT 10
US-10-092-750-1
; Sequence 1, Application US/10092750
; Publication No. US20030032157A1
; GENERAL INFORMATION:
; APPLICANT: Hammond, Philip W.
; APPLICANT: Alpin, Julia
; APPLICANT: Wright, Martin C.
; TITLE OF INVENTION: Polypeptides Interactive with BCL-X1
; FILE REFERENCE: 50036/050002
; CURRENT APPLICATION NUMBER: US/10/092,750
; PRIOR FILING DATE: 2002-03-07
; PRIOR APPLICATION NUMBER: US 60/274,526
; PRIOR FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 253
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 35
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-092-750-1

Query Match      34.1%; Score 47; DB 15; Length 35;
Best Local Similarity 45.5%; Pred. No. 5;
Matches 10; Conservative 3; Mismatches 5; Indels 4; Gaps 1;

QY      2 LWAAQRYGRLRRMSDEFGSF 23
DB      15 IWIAQ----ELRRIGDEFNAY 32

RESULT 11
US-10-092-750-241
; Sequence 241, Application US/10092750
; Publication No. US20030032157A1
; GENERAL INFORMATION:
; APPLICANT: Hammond, Philip W.
; APPLICANT: Alpin, Julia
; APPLICANT: Wright, Martin C.
; TITLE OF INVENTION: Polypeptides Interactive with BCL-X1
; FILE REFERENCE: 50036/050002
; CURRENT APPLICATION NUMBER: US/10/092,750
; PRIOR FILING DATE: 2002-03-07
; PRIOR APPLICATION NUMBER: US 60/274,526
; PRIOR FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 253
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 241
; LENGTH: 138
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-092-750-241

Query Match      34.1%; Score 47; DB 15; Length 138;
Best Local Similarity 45.5%; Pred. No. 20;
Matches 10; Conservative 3; Mismatches 5; Indels 4; Gaps 1;

QY      2 LWAAQRYGRLRRMSDEFGSF 23
DB      86 IWIAQ----ELRRIGDEFNAY 103

RESULT 12
US-10-238-075-1077
; Sequence 1077, Application US/10238075
; Publication No. US20030148324A1
; GENERAL INFORMATION:
; APPLICANT: I.N.S.E.R.M.
; TITLE OF INVENTION: Polynucleotides which are of nature B2/D+ A- and which are iso
; FILE REFERENCE: BLANDINE
; CURRENT APPLICATION NUMBER: US/10/238,075
; CURRENT FILING DATE: 2002-09-10
; PRIOR APPLICATION NUMBER: 0003145
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 1576
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1077
; LENGTH: 682
; TYPE: PRT
; ORGANISM: Escherichia coli
US-10-238-075-1077

Query Match      33.3%; Score 46; DB 12; Length 682;
Best Local Similarity 40.0%; Pred. No. 1.5e+02;
Matches 10; Conservative 5; Mismatches 10; Indels 0; Gaps 0;

QY      2 LWAAQRYGRLRRMSDEFGSKGL 26
DB      610 IWAAQRNGAKVPRVRNGFTSMDIGL 634

RESULT 13
US-09-828-423-3
; Sequence 3, Application US/09828423
; Patent No. US20020099178A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Guegler, Karl J.
; Patterson, Chandra
; TITLE OF INVENTION: GROWTH-ASSOCIATED TRIPSPIN-TYPE
; INHIBITOR HEAVY CHAIN PRECURSOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: Word Perfect 6.1/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/828,423
; FILING DATE: 05-Apr-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/388,774
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Cerrone, Michael C
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0505 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
```

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;
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
;   LENGTH: 946 amino acids
;   TYPE: amino acid
;   STRANDEDNESS: single
;   TOPOLOGY: linear
; IMMEDIATE SOURCE:
;   LIBRARY: GENEBANK
;   CLONE: gi33985
; SEQUENCE DESCRIPTION: SEQ ID NO: 3 :
US-09-828-423-3
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Query Match          33.3%;   Score 46;   DB 9;   Length 946;
Best Local Similarity 30.8%;   Pred. No. 2.le+02;
Matches 8;   Conservative 5;   Mismatches 13;   Indels 0;   Gaps 0;
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QY 1 NLWAAQRYGRLRRMSDEFGSGKGL 26
   :: : | : | : | | | | :
Db 212 DWVIEPQGLRFLHVPDTTEGHFDGV 237
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RESULT 14

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US-10-156-761-11541
; Sequence 11541, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156.761
; PRIOR FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 11541
; LENGTH: 272
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-11541
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Query Match          31.9%;   Score 44;   DB 15;   Length 272;
Best Local Similarity 53.3%;   Pred. No. 1.le+02;
Matches 8;   Conservative 2;   Mismatches 5;   Indels 0;   Gaps 0;
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QY 3 WAAQRYGRLRRMSD 17
   | | : | | | | :
Db 29 WIAAAGAEELRAAD 43
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RESULT 15

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US-09-815-242-5704
; Sequence 5704, Application US/09815242
; Patent No. US20020061569A1
; GENERAL INFORMATION:
; APPLICANT: Haselbeck, Robert
; APPLICANT: Ohlson, Karl L.
; APPLICANT: Zyskind, Judith W.
; APPLICANT: Wall, Daniel
; APPLICANT: Trawick, John D.
; APPLICANT: Carr, Grant J.
; APPLICANT: Yamamoto, Robert T.
; APPLICANT: Xu, H. Howard
; TITLE OF INVENTION: Identification of Essential Genes in
; PROKARYOTES
```

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; FILE REFERENCE: ELITRA.011A
; CURRENT APPLICATION NUMBER: US/09/815.242
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: 60/191,078
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: 60/206,848
; PRIOR FILING DATE: 2000-05-23
; PRIOR APPLICATION NUMBER: 60/207,727
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: 60/242,578
; PRIOR FILING DATE: 2000-10-23
; PRIOR APPLICATION NUMBER: 60/253,625
; PRIOR FILING DATE: 2000-11-27
; PRIOR APPLICATION NUMBER: 60/257,931
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: 60/269,308
; PRIOR FILING DATE: 2001-02-16
; NUMBER OF SEQ ID NOS: 14110
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5704
; LENGTH: 426
; TYPE: PRT
; ORGANISM: Staphylococcus aureus
US-09-815-242-5704

Query Match          31.9%;   Score 44;   DB 9;   Length 426;
Best Local Similarity 36.8%;   Pred. No. 1.8e+02;
Matches 7;   Conservative 4;   Mismatches 8;   Indels 0;   Gaps 0;
```

```
QY 8 YGRLRRMSDEFGSGKGL 26
   : | | | : | | | :
Db 386 FGGSLRRQDENFDGKIKAI 404
```

```
Search completed: September 15, 2003, 17:47:52
Job time : 21.6143 secs
```


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OM protein - protein search, using sw model

Run On: September 15, 2003, 17:18:16 ; Search time 11.7 Seconds
(without alignments)
213.708 Million cell updates/sec

Title: US-09-544-664-1

Perfect score: 138
Sequence: 1 NLWAAQRYGRLRMDSDEFGSKGL 26

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues
Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_76: *
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	138	100.0	204	2 A55671	bad protein - mous
2	54	39.1	946	2 JC5575	inter-alpha-trypsi
3	53	38.4	223	2 D70760	hypothetical prote
4	53	38.4	946	2 S54354	inter-alpha-inhibi
5	52	37.7	370	2 S38185	2-dehydro-3-deoxy-
6	51	37.0	232	2 A42095	floral homeotic pr
7	50	36.2	374	2 C84338	spermidine/putresc
8	50	36.2	516	2 A96753	probable threonine
9	49	35.5	453	2 E83517	conserved hypothet
10	48.5	35.1	134	2 S40376	19 kappa chain - h
11	48.5	35.1	314	2 T02975	annexin p35 - maiz
12	48	34.8	206	2 C36365	transforming prote
13	48	34.8	220	2 F72289	oxidoreductase, so
14	48	34.8	526	2 T08545	threonine synthase
15	47	34.1	597	2 G82308	oxaloacetate decar
16	47	34.1	967	2 F82668	oxoglutarate dehyd
17	47	34.1	5138	2 B96695	hypothetical prote
18	46.5	33.7	314	2 T02961	annexin p33 - maiz
19	46.5	33.7	435	2 A44308	Antho-BFamide prec
20	46.5	33.7	1140	2 T09486	hypothetical prote
21	46	33.3	399	2 T35440	probable polyamine
22	46	33.3	946	1 YHU2	inter-alpha-trypsi
23	46	33.3	1164	2 T24806	hypothetical prote
24	46	33.3	1378	2 A81993	DNA-directed RNA p
25	45.5	33.0	261	2 G69510	conserved hypothet
26	45.5	33.0	287	2 S43852	neuropeptide pol-R
27	45.5	33.0	334	2 A39172	Antho-BFamide neur
28	45.5	33.0	562	2 C71473	hypothetical prote
29	45.5	33.0	905	2 G63314	NADH dehydrogenase

ALIGNMENTS

RESULT 1

A55671
bad protein - mouse
C:Species: Mus musculus (house mouse)
C>Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999
C/Accession: A55671
R:Yang, E.; Zha, J.; Jockel, J.; Boise, L.H.; Thompson, C.B.; Korsmeyer, S.J.
Cell 80, 285-291, 1995
A>Title: Bad, a heterodimeric partner for Bcl-x-L and Bcl-2, displaces Bax and promot
A:Reference number: A55671; MUID:95136361; PMID:7834748
A:Accession: A55671
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-204 <YAN>
A:Cross-references: GB:L37296; NID:9639778; PIDN:AAA64465.1; PID:G639779
C:Keywords: heterodimer

Query Match 100.0%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 7e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRLRMDSDEFGSKGL 26
DB 140 NLWAAQRYGRLRMDSDEFGSKGL 165

RESULT 2

JC5575
inter-alpha-trypsin inhibitor heavy chain 2 - golden hamster
C:Species: Mesocricetus auratus (golden hamster)
C>Date: 23-Sep-1997 #sequence_revision 23-Sep-1997 #text_change 20-Jun-2000
C/Accession: JC5575; PC4485
R:Nakatani, T.; Suzuki, Y.; Yamamoto, T.; Sinohara, H.
J. Biochem. 122, 71-82, 1997
A>Title: Molecular cloning and sequencing of cDNAs encoding three heavy-chain precu
sin inhibitor heavy chain family.
A:Reference number: JC5574; MUID:97420688; PMID:9276673
A:Accession: JC5575
A:Molecule type: mRNA
A:Residues: 1-946 <NAK>
A:Cross-references: DDBJ:D99286; NID:G1694689; PIDN:BAA13939.1; PID:G1694690
A:Experimental source: liver
A:Accession: PC4485
A:Molecule type: protein
A:Residues: 55-64;140-146;151-156;424-447;500-528;577-605 <NA2>
C:Comment: In the plasma three inter-alpha-trypsin inhibitor heavy chains 1, 2 and 3
that the complexes play important role for pancreatic cancer.
C:Superfamily: inter-alpha-trypsin inhibitor complex component II
P:261-264,717-916/Disulfide bonds: #status predicted

Query Match 39.1%; Score 54; DB 2; Length 946;
Best Local Similarity 34.6%; Pred. No. 8.8;

C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001

C:Accession: A96753
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alon
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar,
ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000

A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim,
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marzia
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallo
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:title: Sequence and analysis of chromosome 1 of the plant *Arabidopsis*.
A:Reference number: A86141; MUID:21016719; PMID:11130712
A:Accession: A96753
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-516 <STO>
A:Cross-references: GB:AE005173; NID:g5903070; PIDN:AAD55628.1; GSPDB:GN00141
C:Genetics:
A:Gene: F3N23.1
A:Map position: 1

Query Match 36.2%; Score 50; DB 2; Length 516;
Best Local Similarity 35.3%; Pred. NO.18;
Matches 12; Conservative 7; Mismatches 7; Indels 8; Gaps 1;

QY 1 NLWAAORYGRELRLMSD-----EFEGSFKGL 26
|||:||||:||||
Db 163 NLFWAFRFKQVLQNDLWKHCGISHTGSFKDL 196

RESULT 9
E83517
conserved hypothetical protein PA1031 [imported] - *Pseudomonas aeruginosa* (strain PAO)
C:Species: *Pseudomonas aeruginosa*
C>Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: E83517
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.;
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; L
Lory, S.; Olson, M.V.
Nature 406, 949-964, 2000

A:title: Complete genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic pa
A:Reference number: A82950; MUID:20437337; PMID:10984043
A:Accession: E83517
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-453 <STO>
A:Cross-references: GB:AE004535; GB:AE004091; NID:g9946936; PIDN:AG04420.1; GSPDB:GN
A:Experimental source: strain PA01
C:Genetics:
A:Gene: PA1031

Query Match 35.5%; Score 49; DB 2; Length 453;
Best Local Similarity 55.6%; Pred. NO.23;
Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;

QY 3 WAAQRYGR--ELRRMSDE 18
|||:||| |||||:|
Db 65 WASEROGREELRLASE 82

RESULT 10
S40376
Ig kappa chain - human
C:Species: Homo sapiens (man)
C>Date: 06-Mar-1994 #sequence_revision 26-May-1995 #text_change 21-Jan-2000
C:Accession: S40376
R:Klein, R.; Jeenichen, R.; Zachau, H.G.
Eur. J. Immunol. 23, 3248-3271, 1993

A:title: Expressed human immunoglobulin chi genes and their hypermutation.
A:Reference number: S40312; MUID:94080891; PMID:8258341
A:Accession: S40376
A>Status: preliminary; translation not shown

Qy.

```
Db      172 NLFWAERFGQFLGNDLWVHCIGISHTGSKDL 205
|||||
RESULT 15
G82308
oxaloacetate decarboxylase, alpha chain VC0550 [similarity] - Vibrio cholerae (strain N1
C;Species: Vibrio cholerae
C;Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
C;Accession: G82308
R;Heidelberger, J.F.; Eissen, J.A.; Nelson, W.C.; Clayton, R.A.; Winn, M.L.; Dodson, R.J.;
chardson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragoi, I.; Sellers, R
l, R.R.; Melanos, J.J.; Venter, J.C.; Fraser, C.M.
Nature 406, 477-483, 2000
A;Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.
A;Reference number: A82035; MUID:20406833; PMID:10952301
A;Accession: G82308
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-597 <HEI>
A;Cross-references: GB:AE004141; GB:AE003852; NID:99554976; PIDN:AAF93718.1; GSPDB:GN001
A;Experimental source: serogroup O1; strain N16961; biotype El Tor
C;Genetics:
A;Gene: VC0550
A;Map position: 1
C;Superfamily: Klebsiella pneumoniae oxaloacetate decarboxylase alpha chain; lipoyl/biot
Query Match      34.1%; Score 47; DB 2; Length 597;
Best Local Similarity 47.4%; Pred. No. 60;
Matches 9; Conservative 4; Mismatches 6; Indels 0; Gaps 0;
QY      8 YGRELRMSDEFESEKGL 26
Db      272 YFEVRKKYAKFEGQLKV 290
|||||
Search completed: September 15, 2003, 17:27:00
Job time : 12.7 secs
```

GenCore version 5.1.6
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OM protein - protein search, using sw model,
Run on: September 15, 2003, 17:16:55; Search time 6.12857 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-1
Perfect score: 138
Sequence: 1 NLWAAQRYGRLRMSDFEGSKGL 26

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	138	100.0	204	1 BAD_MOUSE	Q61337 mus musculus
2	138	100.0	205	1 BAD_RAT	Q35147 rattus norv
3	114	82.6	168	1 BAD_HUMAN	Q92934 homo sapien
4	54	39.1	946	1 ITH2_MESAU	P97279 mesocricetu
5	53	38.4	946	1 ITH2_MOUSE	Q61703 mus musculu
6	52	37.7	370	1 AROG_YEAST	P32449 saccharomyc
7	51	37.0	232	1 AP3_ARATH	P35632 arabidopsis
8	50	36.2	851	1 CE05_MOUSE	Q8K2H3 mus musculu
9	49.5	35.9	506	1 MATK_LEDPA	Q62992 ledum palus
10	49.5	35.9	506	1 MATK_RHOFR	Q62984 rhododendro
11	49.5	35.9	506	1 MATK_RHOTS	Q62991 rhododendro
12	49	35.5	453	1 RMUC_PSEAE	Q914U3 pseudomonas
13	48	34.8	205	1 RAS3_RHIRA	P22280 rhizomucor
14	48	34.8	220	1 6PGL_THEMA	Q9X0N8 thermotoga
15	48	34.8	519	1 THRC_SOLTU	Q9MT28 solanum tub
16	48	34.8	526	1 THRC_ARATH	Q9S7B5 arabidopsis
17	47	34.1	198	1 BIM_HUMAN	Q43521 homo sapien
18	46.5	33.7	429	1 FMR2_ANTEL	Q16994 anthopleura
19	46.5	33.7	435	1 FMR1_ANTEL	P10419 anthopleura
20	46	33.3	946	1 ITH2_HUMAN	P19823 homo sapien
21	46	33.3	1378	1 RPOB_CAMJE	Q46124 campylobact
22	45.5	33.0	287	1 PRFA_POLPE	P21259 polyorchis
23	45.5	33.0	334	1 FMRA_CALPA	Q01133 calliactis
24	45.5	33.0	507	1 MATK_LOIPR	Q47169 loiseleuria
25	45	32.6	328	1 SNF4_KLULA	Q9P869 kluyveromyc
26	45	32.6	590	1 DCOA_SALTY	Q03030 salmonella
27	45	32.6	595	1 HTOA_KLEPN	P13187 klebsiella
28	45	32.6	653	1 ITH2_HUMAN	Q13049 homo sapien
29	45	32.6	865	1 ENV_SIVAG	P05886 simian immu
30	45	32.6	1535	1 LML1_CAEEL	Q18923 caenorhabdi
31	44.5	32.2	907	1 NUOG_ECOLI	P33602 escherichia
32	44.5	32.2	907	1 NUOG_SALTY	P33900 salmonella
33	44	31.9	196	1 BIM_MOUSE	Q54918 mus musculu

RESULT 1

ID	BAD_MOUSE	STANDARD;	PRT;	204 AA.
AC	Q61337;			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	28-FEB-2003 (Rel. 41, Last annotation update)			
DE	Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component			
DE	6) (Bcl-XL/Bcl-2 associated death promoter).			
GN	BAD OR BOC6.			
OS	Mus musculus (Mouse).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;			
OX	NCBI_TaxID=10090;			
[1]				
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Brain, and Thymus;			
RX	MEDLINE=95136361; PubMed=7834748;			
RA	Yang E., Zhu J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;			
RT	"Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and			
RT	promotes cell death.";			
RL	Cell 80:285-291(1995).			
[2]				
RP	PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.			
RX	MEDLINE=98022383; PubMed=9381178;			
RA	Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;			
RT	"Interleukin-3-induced phosphorylation of BAD through the protein			
RT	kinase Akt.";			
RL	Science 278:687-689(1997).			
[3]				
RP	MUTAGENESIS OF SERINE RESIDUES.			
RX	MEDLINE=20403302; PubMed=10949026;			
RA	Datta S.R., Katsav A., Hu L., Petros A., Fesik S.W., Yaffe M.B.;			
RT	Greenberg M.E.;			
RT	"14-3-3 proteins and survival kinases cooperate to inactivate BAD by			
RT	BH3 domain phosphorylation.";			
RL	Mol. Cell 6:41-51(2000).			
CC	-I- FUNCTION: Promotes cell death. Successfully competes for the			
CC	binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level			
CC	of heterodimerization of these proteins with BAX. Can reverse the			
CC	death repressor activity of Bcl-x(L), but not that of Bcl-2.			
CC	Appears to act as a link between growth factor receptor signaling			
CC	and the apoptotic pathways.			
CC	-I- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-			
CC	x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).			
CC	The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.			
CC	-I- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon			
CC	phosphorylation, locates to the cytoplasm.			
CC	-I- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND			
CC	BAX for their pro-apoptotic activity and for their interaction			
CC	with anti-apoptotic members of the Bcl-2 family.			
CC	-I- PTM: Phosphorylated on Ser-112 in response to survival stimuli.			
CC	Subsequent phosphorylation on Ser-136 promotes heterodimerization			
CC	with 14-3-3 proteins. This interaction then facilitates the			
CC	phosphorylation at Ser-155, a site within the BH3 domain, leading			
CC	to the release of Bcl-x(L) and the promotion of cell survival.			

ALIGNMENTS

34	44	31.9	196	1 BIM_RAT	Q88498 rattus norv
35	44	31.9	262	1 END8_ECO57	Q8X9C6 escherichia
36	44	31.9	262	1 END8_ECOLI	P50465 escherichia
37	44	31.9	262	1 END8_SALTY	Q8Z832 salmonella
38	44	31.9	262	1 END8_SALTY	Q8Z832 salmonella
39	44	31.9	768	1 ENV_SIVAG	P27757 simian immu
40	44	31.9	877	1 ENV_SIVAG	P27757 simian immu
41	44	31.9	915	1 CE05_HUMAN	Q9NYF5 homo sapien
42	44	31.9	5596	1 MDN1_HUMAN	Q9NU22 homo sapien
43	44	31.9	8797	1 SNE1_HUMAN	Q8NF91 homo sapien
44	43.5	31.5	217	1 UREF_SYNY3	P73327 synechocyst
45	43.5	31.5	1014	1 UVRA_STRCO	Q9Z507 streptomyce

RA Hamner S., Arumae U., Yu L.-Y., Sun Y.-F., Saarma M., Lindholm D.;
RT "Functional characterization of two splice variants of rat BAD and
TT their interaction with Bcl-w in sympathetic neurons.";
RL Mol. Cell. Neurosci. 17:97-106(2001).
CC -!- FUNCTION: Promotes cell death. Successfully competes for the
CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
CC of heterodimerization of these proteins with BAX. Can reverse the
CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
CC similarity). Appears to act as a link between growth factor
CC receptor signaling and the apoptotic pathways.
CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
CC x(L), Bcl-2 and Bcl-w. Also binds protein SL100A10. The Ser-
CC 113/Ser-137 phosphorylated form binds 14-3-3 proteins.
CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
CC phosphorylation, locates to the cytoplasm (By similarity).
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=Alpha;
CC IsoId=Q35147-1; Sequence=Displayed;
CC Name=Beta;
CC IsoId=Q35147-2; Sequence=VSP_000534;
CC -!- TISSUE SPECIFICITY: Expressed in all tissues tested, including
CC brain, liver, spleen and heart. In the brain, restricted to
CC epithelial cells of the choroid plexus. Isoform alpha is the more
CC abundant form.
CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
CC BAX for their pro-apoptotic activity and for their interaction
CC with anti-apoptotic members of the Bcl-2 family.
CC -!- PTM: Phosphorylated on Ser-113 in response to survival stimuli.
CC subsequent phosphorylation on Ser-137 promotes heterodimerization
CC with 14-3-3 proteins. This interaction then facilitates the
CC phosphorylation at Ser-156, a site within the BH3 domain, leading
CC to the release of Bcl-x(L) and the promotion of cell survival.
CC Ser-137 is the major site of AKT/PKB phosphorylation, Ser-156 the
CC major site of protein kinase A (CAPK) phosphorylation (By
CC similarity).
CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: AF003523; AAC53374.1; -
DR EMBL: AF031227; AAC15100.1; -
DR ENBL: AF279910; AAF9427.1; -
DR ENBL: AF279911; AAF9428.1; -
DR HSSP: Q29334; IG5J.
DR InterPro: IPR000712; BCl2_BH.
DR PROSITE: PS01259; BH3; FALSE NEG.
KW Apoptosis; Phosphorylation; Alternative splicing.
FT FT MOD_RES 148 162 BH3 PHOSPHORYLATION (BY PKA AND PKB)
FT FT MOD_RES 137 137 (BY SIMILARITY).
FT FT MOD_RES 156 156 PHOSPHORYLATION (BY PKA AND PKB)
FT FT MOD_RES 156 156 (BY SIMILARITY).
FT FT MOD_RES 156 156 PHOSPHORYLATION (BY PKA AND PKB)
FT FT VARSPLIC 166 205 (LPKRSAGTATQRKSASWTLIISWDNRLKGSGTPSQ
FT -> BELTVSEFLPVFAIMGWLIVSFQSFPHTLPPTPP
FT EVAMFLPYRVALRRLLC (in isoform Beta).
FT FT FTId=VSP_000534.
FT S->> NO EFFECT ON HETERODIMERIZATION
FT WITH 14-3-3 PROTEINS.
FT S->> NO HETERODIMERIZATION WITH 14-3-3
FT PROTEINS. NO EFFECT ON HETERODIMERIZATION
FT WITH BCL2 NOR WITH PROTEIN P11.
FT SDAGGR -> ERRGR (IN REF. 1).
FT CONFLICT 29 34
FT SOURCE 205 AA; 22228 MW; 7AFAY1DAE9CA81 CRC64;

Query Match 100.0%; Score 138; DB 1; Length 205;
 Best Local Similarity 100.0%; Pred. No. 1.6e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRELARMDSDEFGSKGL 26
 Db 141 NLWAAQRYGRELARMDSDEFGSKGL 166

RESULT 3

BAD_HUMAN STANDARD; PRT: 168 AA.
 ID BAD_HUMAN Q92934; O14803;
 AC Q92934; O14803;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 15-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-
 XL/Bcl-2 associated death promoter) (BCL2-like 8 protein).
 GN BAD OR BCL2L8.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 OX NCBI_TaxID=9606;
 [1]
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Yin D.X., Li Z., Huang B., Chen S., Zhou H.;
 RT "A human protein that interacts with Bcl-2 and have homology to mouse
 RT BAD.";
 RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.
 RX MEDLINE=97083574; PubMed=8929532;
 RA Wang H.-G., Rapp U.R., Reed J.C.;
 RT "Bcl-2 targets the protein kinase Raf-1 to mitochondria.";
 RL Cell 87:629-638(1996).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Takayama S., Reed J.C.;
 RL Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A., AND DIMERIZATION.
 RC TISSUE=Bone marrow;
 RX MEDLINE=98049554; PubMed=9388232;
 RA Otilie S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,
 RA Chang S., Weeks S., Fritz L.C., Oltersdorf T.;
 RT "Dimerization properties of human BAD.";
 RL J. Biol. Chem. 272:30866-30872(1997).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lung;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg K.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Atschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Vallalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kerteman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Touchman J.W., Green E.D., Dickson M.C.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalhus D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [6]

RP STRUCTURE BY NMR OF 103-127.
 RX MEDLINE=21073561; PubMed=11206074;
 RA Petros A.M., Nettekheim D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
 RA Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
 RA Resik S.W.;
 RT "Rationale for Bcl-xL/Bad peptide complex formation from structure,
 RT mutagenesis, and biophysical studies.";
 RL Protein Sci. 9:2528-2534(2000).
 CC -!- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 CC similarity). Appears to act as a link between growth factor
 CC receptor signaling and the apoptotic pathways.
 CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 CC The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
 CC similarity).
 CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm.
 CC -!- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
 CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -!- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-99 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-118, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-99 is the major site of AKT/PKB phosphorylation, Ser-118 the
 CC major site of protein kinase A (CAPK) phosphorylation (by
 CC similarity).
 CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
 CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -!- CAUTION: Ref.1 sequence differs from that shown due to frameshifts
 CC in position 64 and 91.
 CC -----
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 CC -----
 CC EMBL; U66879; AAB36516.1; ALT_FRAME.
 CC EMBL; AF021792; AAB72092.1; -.
 CC EMBL; AF031523; AAB88124.1; -.
 CC EMBL; BC001901; AAB01901.1; -.
 CC PDB; 1G5J; 07-FEB-01.
 CC Genew; HGNC:936; BND.
 CC MIM; 603167; -.
 CC GO; GO:0005737; C:cytoplasm; NAS.
 CC GO; GO:0005741; C:mitochondrial outer membrane; NAS.
 CC GO; GO:0005515; F:protein binding activity; NAS.
 CC GO; GO:0008632; P:apoptotic program; TAS.
 CC GO; GO:0006917; P:induction of apoptosis; NAS.
 CC InterPro; IPR000712; Bcl2_BH.
 CC PROSITE; PS01259; BH3; FALSE_NEG.
 CC Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
 CC DOMAIN 110 124
 CC MOD_RES 75 75
 CC PHOSPHORYLATION (BY PKA AND PKB) (BY
 CC SIMILARITY).
 CC MOD_RES 99 99
 CC PHOSPHORYLATION (BY PKA AND PKB) (BY
 CC SIMILARITY).
 CC MOD_RES 118 118
 CC PHOSPHORYLATION (BY PKA AND PKB) (BY
 CC SIMILARITY).
 CC VARIANT 107 107
 CC A -> S (in dbSNP:3729933).
 CC /FTID=VAR_015380.
 CC HELIX 106 121
 CC SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
 SQ

Query Match 82.6%; Score 114; DB 1; Length 168;

[illegible]

RN [2] SEQUENCE FROM N.A.
 RP STRAIN=cv. Landsberg erecta;
 RC MEDLINE=95036018; PubMed=7948893;
 RA Okamoto H., Yano A., Shiraishi H., Okada K., Shimura Y.;
 RT "Genetic complementation of a floral homeotic mutation, *apetala3*,
 RT with an Arabidopsis thaliana gene homologous to *DEFICIENS* of
 RT Antirrhinum majus.";
 RL Plant Mol. Biol. 26:465-472(1994).
 RN [3] SEQUENCE FROM N.A., AND VARIANTS.
 RP STRAIN=cv. Bla-1, cv. Bretagny, cv. Bs-1, cv. Bu-0, cv. Bu-2,
 RC cv. Chi-1, cv. Co-1, cv. Columbia, cv. Corsacalia-1, cv. Cvi-0,
 RC cv. Gr-3, cv. J1-1, cv. Kas-1, cv. Kent, cv. Landsberg erecta,
 RC cv. Li-3, cv. Li-8, and cv. Lisse;
 RX MEDLINE=99126449; PubMed=9927474;
 RA Purganan M.D., Suddith J.I.;
 RT "Molecular population genetics of floral homeotic loci: departures
 RT from the equilibrium-neutral model at the *APETALA3* and *PISTILLATA*
 RT genes of Arabidopsis thaliana.";
 RL Genetics 151:839-848(1999).
 RN [4] SEQUENCE FROM N.A.
 RP STRAIN=cv. Columbia;
 RC MEDLINE=21016720; PubMed=11130713;
 RX Salanoubat M., Lemcke K., Rieger M., Ansoerge W., Unseld M.,
 RA Fartmann B., Valle G., Bloeker H., Perez-Alonso M., Obermaier B.,
 RA Delseny M., Boutry M., Griuell L.A., Mache R., Puigdomenech P.,
 RA De Simone V., Choisine N., Artiguenave F., Robert C., Brottier P.,
 RA Wincker P., Cattolico L., Weissbach J., Saurin W., Quetier F.,
 RA Schaefer M., Mueller-Auer S., Gabel C., Fuchs M., Benes V.,
 RA Wurmacher E., Drzonek H., Erle H., Jordan N., Bangert S.,
 RA Wiedelmann R., Kranz H., Voss H., Holland R., Brandt P., Nyakatura G.,
 RA Vezzi A., D'Angelo M., Pallavicini A., Roppo S., Simonati B.,
 RA Conrad A., Hornischer K., Kauer G., Loehner T.-H., Nordstiek G.,
 RA Reichelt J., Collado C., Schoen O., Barques M., Terol J., Climent J.,
 RA Navarro P., Collado C., Perez-Perez A., Ottenwaelder B., Duchemin D.,
 RA Cooke R., Landie M., Berger-Lilauro C., Purnelle B., Masuy D.,
 RA de Haan M., Maarse A.C., Alcaraz J.-P., Cottet A., Casacuberta E.,
 RA Monfort A., Argivon A., Flores M., Liguori R., Vitale D.,
 RA Mannhaupt G., Haase D., Schoof H., Rudd S., Zaccaria P., Meves H.-W.,
 RA Mayer K.F.X., Kaul S., Town C.D., Koo H.B., Tallon L.J., Jenkins J.,
 RA Rooney T., Rizzo M., Walts A., Utterback T., Fujii C.Y., Shea T.P.,
 RA Cressy T.H., Haas B., Mafti R., Wu D., Peterson J., Van Aken S.,
 RA Pai G., Miltscher J., Sellers P., Gill J.E., Feldblyum T.V.,
 RA Preuss D., Lin X., Nierman W.C., Salzberg S.L., White O., Venter J.C.,
 RA Fraser C.M., Kaneko T., Nakamura Y., Sato S., Kato T., Asamizu E.,
 RA Sasamoto S., Kimura T., Idesawa K., Kawashima K., Kishida Y.,
 RA Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
 RA Nakayama S., Nakazaki N., Shinpo S., Takeuchi C., Wada T.,
 RA Watanabe A., Yamada M., Yasuda M., Tabata S.;
 RT "Sequence and analysis of chromosome 3 of the plant Arabidopsis
 RT thaliana.";
 RL Nature 408:820-822(2000).
 RN [5] SEQUENCE FROM N.A.
 RP Brover V., Troukhan M., Alexandrov N., Lu Y.-P., Flavell R.,
 RA Feldmann K.;
 RT "Full-length cDNA from Arabidopsis thaliana.";
 RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
 RN [6] SEQUENCE FROM N.A.
 RP STRAIN=cv. Columbia;
 RA Shinozaki K., Davis R.W., Ecker J.R., Theologis A.;
 RT "RIKEN Arabidopsis full length cDNA clones (RAFLs) sequenced by the
 RT SSP consortium (Salk/Stanford/PGECC).";
 RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
 RN [7] SEQUENCE OF 36-128 FROM N.A.
 RP STRAIN=cv. Columbia;
 RC MEDLINE=99311297; PubMed=10382288;
 RX Brunel D., Froger N., Pelletier G.;
 RA "Development of amplified consensus genetic markers (ACGM) in Brassica

RT napus from Arabidopsis thaliana sequences of known biological
 RT function.";
 RL Genome 42:387-402(1999).
 RN [8] FUNCTION.
 RP PubMed=8565821;
 RA Krizek B.A., Meyerowitz E.M.;
 RT "The Arabidopsis homeotic genes *APETALA3* and *PISTILLATA* are sufficient
 RT to provide the B class organ identity function.";
 RL Development 122:11-22(1996).
 RN [9] CHARACTERIZATION.
 RP PubMed=8643482;
 RA Riechmann J.L., Krizek B.A., Meyerowitz E.M.;
 RT "Dimerization specificity of Arabidopsis MADS domain homeotic proteins
 RT *APETALA1*, *APETALA3*, *PISTILLATA*, and *AGAMOUS*.";
 RL Proc. Natl. Acad. Sci. U.S.A. 93:4793-4798(1996).
 RN [10] GENETIC REGULATION.
 RP PubMed=11283333;
 RX Ng M., Yano-sky M.F.;
 RA "Activation of the Arabidopsis B class homeotic genes by *APETALA1*.";
 RL Plant Cell 13:739-753(2001).
 RN [11] CHARACTERIZATION.
 RP PubMed=11206550;
 RX Honma T., Goto K.;
 RA "Complexes of MADS-box proteins are sufficient to convert leaves into
 RT floral organs.";
 RL Nature 409:525-529(2001).
 CC -!- FUNCTION: Probable transcription factor involved in the genetic
 CC control of flower development. Is required for normal development
 CC of petals and stamens in the wild-type flower. Forms an
 CC heterodimer with *PISTILLATA* that is required for autoregulation of
 CC both *AP3* and *PI* genes. *AP3/PI* heterodimer interacts with *APETALA1*
 CC or *SEPALLATA3* to form a ternary complex that could be responsible
 CC for the regulation of the genes involved in the flower
 CC development.
 CC -!- SUBUNIT: Forms an heterodimer with *PISTILLATA*, capable of binding
 CC to CARG-box sequences. *AP3/PI* heterodimer binds *AP1* or *SEP3* to
 CC form complexes.
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- TISSUE SPECIFICITY: Expressed in petals and stamens.
 CC -!- INDUCTION: Positively regulated by the meristem identity proteins
 CC *APETALA1* and *LEAFY* with the cooperation of *UFO*.
 CC -!- MISCELLANEOUS: Mutations in *AP3* cause transformation of petals
 CC into sepals and stamens into carpels.
 CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
 CC FACTORS.
 CC -!- SIMILARITY: Contains 1 K-box dimerization domain.
 CC -----
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 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
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 CC -----
 CC EMBL: M66357; AAA32740.1; -;
 CC EMBL: D21125; BAA04665.1; -;
 CC EMBL: AF115798; AAD51887.1; -;
 CC EMBL: AF115799; AAD51888.1; -;
 CC EMBL: AF115800; AAD51889.1; -;
 CC EMBL: AF115801; AAD51890.1; -;
 CC EMBL: AF115802; AAD51891.1; -;
 CC EMBL: AF115803; AAD51892.1; -;
 CC EMBL: AF115804; AAD51893.1; -;
 CC EMBL: AF115805; AAD51894.1; -;
 CC EMBL: AF115806; AAD51895.1; -;
 CC EMBL: AF115807; AAD51896.1; -;
 CC EMBL: AF115808; AAD51897.1; -;
 CC EMBL: AF115809; AAD51898.1; -;
 CC EMBL: AF115809; AAD51898.1; -;

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DR EMBL; AF115810; AAD51899.1; -
DR EMBL; AF115811; AAD51900.1; -
DR EMBL; AF115812; AAD51901.1; -
DR EMBL; AF115813; AAD51902.1; -
DR EMBL; AF115814; AAD51903.1; -
DR EMBL; AL132971; CAB81799.1; -
DR EMBL; AY087369; AAM64919.1; -
DR EMBL; AY070397; AAL49893.1; -
DR EMBL; AY142590; AAN13159.1; -
DR EMBL; AF056541; ABD41557.1; -
DR F1R; A42095; A42095.
DR HSP; P11746; IJNM.
DR TRANSFAC; T01776; -.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADS_DOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS0066; MADS_BOX_2; 1.
KW Flowering; Transcription regulation; Activator; Developmental protein;
KW Nuclear protein; DNA-binding; Coiled coil; Polymorphism.
FT DOMAIN 3 57 MADS.
FT DOMAIN 93 165 K-BOX.
FT DOMAIN 75 164 COILED COIL (POTENTIAL).
FT VARIANT 31 31 K -> R (in strain cv. Lisae).
FT VARIANT 47 47 M -> T (in strain cv. Bretagne).
FT VARIANT 61 61 N -> D (in strain cv. Corsacalla-1).
FT VARIANT 73 73 T -> S (in strain cv. Li-8).
FT VARIANT 109 109 L -> V (in strain cv. Kas-1).
FT VARIANT 115 115 E -> K (in strains cv. Chi-1 and cv. Gr-3).
Query Match 37.0%; Score 51; DB 1; Length 232;
Best Local Similarity 44.4%; Pred. No. 2.1;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEPESFK 24
||| :||||| ||| :||
Db 107 QRLGCEGLDLDIQELRRLEDEMENTFK 133

RESULT 8
CE05_MOUSE STANDARD; PRT; 851 AA.
AC Q8K2H3;
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Protein C5orf5 homolog.
GN C5ORF5.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=FVB/N;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.F., Rubin G.M., Hong L.,
RA Stappleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,

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RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Generch A., Schein J.E., Jones S.J.M., Marra M.A.:
RT "Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- SIMILARITY: Belongs to the FAM13 family.
CC -!- SIMILARITY: Contains 1 Rho-GAP domain.
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CC
CC EMBL; BC031465; AAH31465.1; -.
CC InterPro; IPR000198; RhoGAP.
CC Pfam; PF00620; RhoGAP; 1.
CC SMART; SMO0324; RhoGAP; 1.
CC PROSITE; PS02338; RHO-GAP; 1.
KW GTPase activation. 212 RHO-GAP.
FT DOMAIN 123 256 GLU-RICH.
FT DOMAIN 189 256
SQ SEQUENCE 851 AA; 97054 MW; C2B26669FB6DB2CE CRC64;
Query Match 36.2%; Score 50; DB 1; Length 851;
Best Local Similarity 45.5%; Pred. No. 12;
Matches 10; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 2 LMAAQRYGRELRRMSDEPESGF 23
||| :||||| ||| :||
Db 784 LKWARAEKKKKLREFEAF 805

RESULT 9
MATK_LEDPA STANDARD; PRT; 506 AA.
AC O62992;
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Maturase K (intron maturase).
GN MATK.
OS Ledum palustre (Wild rosemary).
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75583;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of sectional relationships in the genus
RT Rhododendron (Ericaceae) based on matk sequences.";
RL Shokubutsu Kenkyu Zasshi 73:143-154(1998).
CC -!- FUNCTION: Probably assists in splicing chloroplast group II
introns (By similarity).
CC -!- SIMILARITY: BELONGS TO THE INTRON MATURASE FAMILY 2. MATK
SUBFAMILY.
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AB012751; BAA25872.1; -.

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RESULT 11	
MATK_RHOTS	
ID	MATK_RHOTS
AC	002991;
DT	28-FEB-2003 (Rel. 41, Created)
	STANDARD;
	PRT; 506 AA.

RL Natur

Match	Conservative	Mismatches	Indels	Gaps
Matches	11	5	7	7

CC -!- FUNCTION: Involved in DNA recombination (By similarity).
 CC -!- SIMILARITY: BELONGS TO THE RMCU FAMILY.
 CC -----
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 CC -----
 DR EMBL; A5004535; AAC04420.1; -.
 DR PIR; E83517; E83517.
 DR InterPro; IPR003798; DUF195.
 DR Pfam; PF02646; RmcU; 1.
 KW DNA recombination; Coiled coil; Complete proteome.
 FT DOMAIN 16 201 COILED COIL (POTENTIAL).
 SQ SEQUENCE 453 AA; 51539 MW; 1E7EA97E82EC5E4B CRC64;
 Query Match 35.5%; Score 49; DB 1; Length 453;
 Best Local Similarity 55.6%; Pred. No. 8.6;
 Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;
 QY 3 WAAQYGR--ELREXSD 18
 DB 65 WASERQGREELRLASE 82
 RESULT 13
 ID_RAS3_RHRA STANDARD; PRT; 205 AA.
 AC P22280;
 DT 01-AUG-1991 (Rel. 19, Created)
 DT 01-AUG-1991 (Rel. 19, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Ras-like protein 3.
 GN RAS3.
 OS Rhizomucor racemosus (Mucor circinelloides f. lusitanicus).
 CC Eukaryota; Fungi; Zygomycota; Zygomycetes; Mucorales; Mucoraceae;
 CC Mucor.
 CC NCBI_TaxID=4841;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 1216B;
 RX MEDLINE=91061774; PubMed=1701021;
 RA Casale W.L., McConnell D.G., Wang S.-Y., Lee Y.-J., Linz J.E.;
 RT "Expression of a gene family in the dimorphic fungus Mucor racemosus
 RT which exhibits striking similarity to human ras genes";
 RL Mol. Cell. Biol. 10:6654-6663(1990).
 CC -!- ENZYME REGULATION: ALTERNATE BETWEEN AN INACTIVE FORM BOUND TO GDP
 CC AND AN ACTIVE FORM BOUND TO GTP. ACTIVATED BY A GUANINE
 CC NUCLEOTIDE-EXCHANGE FACTOR (GEF) AND INACTIVATED BY A GTPASE-
 CC ACTIVATING PROTEIN (GAP).
 CC -!- SUBCELLULAR LOCATION: Plasma membrane.
 CC -!- DEVELOPMENTAL STAGE: IN SPORULATING MYCELIUM AND MUCH LESS IN
 CC GERMLING AND YEAST.
 CC -!- SIMILARITY: BELONGS TO THE SMALL GTPASE SUPERFAMILY. RAS FAMILY.
 CC -----
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 CC -----
 DR EMBL; M53177; AAA83379.1; -.
 DR PIR; C36365; C36365.
 DR HSSP; P01112; 1PL1.
 DR InterPro; IPR003577; GTPase_Ras.
 DR InterPro; IPR001806; Ras_trnsmrg.
 DR Pfam; PF00071; ras; 1.

DR PRINTS; PR00449; RASTRNSFRMG.
 DR SMART; SM00173; RAS; 1.
 DR TIGRFAMS; TIGR00231; small_GTP; 1.
 KW GTP-binding; Prenylation; Lipoprotein.
 FT NP_BIND 16 23 GTP (BY SIMILARITY).
 FT NP_BIND 63 67 GTP (BY SIMILARITY).
 FT NP_BIND 122 125 GTP (BY SIMILARITY).
 FT DOMAIN 38 46 EFFECTOR REGION (PROBABLE).
 FT LIPID 202 202 FARNESYL (BY SIMILARITY).
 SQ SEQUENCE 205 AA; 23408 MW; D8F086456F090F50 CRC64;
 Query Match 34.8%; Score 48; DB 1; Length 205;
 Best Local Similarity 62.5%; Pred. No. 5.1;
 Matches 10; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
 QY 10 RELRMSDEFEFGSKG 25
 DB 168 REIRMKKEGERSKG 183
 RESULT 14
 ID_6PGL_THEME STANDARD; PRT; 220 AA.
 AC O3X0N8;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 6-phosphogluconolactonase (EC 3.1.1.31) (6PGL).
 GN PGL OR DEVB OR TM1154.
 OS Thermotoga maritima.
 CC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
 CC NCBI_TaxID=2336;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MSB8 / DSM 3109;
 RX MEDLINE=99287316; PubMed=10360571;
 RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
 RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
 RA McDonald L., Utterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
 RA Stewart A.N., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
 RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
 RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
 RT "Evidence for lateral gene transfer between Archaea and Bacteria from
 RT genome sequence of Thermotoga maritima";
 RL Nature 393:323-329(1999).
 CC -!- FUNCTION: HYDROLYSIS OF 6-PHOSPHOGLUCONOLACTONE TO 6-
 CC PHOSPHOGLUCONATE.
 CC -!- CATALYTIC ACTIVITY: 6-phospho-D-glucono-1,5-lactone + H(2)O = 6-
 CC phospho-D-gluconate.
 CC -!- PATHWAY: Pentose phosphate pathway; second step.
 CC -!- SIMILARITY: BELONGS TO THE GLUCOSAMINE/GALACTOSAMINE-6-PHOSPHATE
 CC ISOMERASE FAMILY. 6-PHOSPHOGLUCONOLACTONASE SUBFAMILY.
 CC -----
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 CC -----
 DR EMBL; AE001772; AAD36230.1; -.
 DR PIR; F72289; F72289.
 DR TIGR; TM1154; -.
 DR InterPro; IPR006148; Gluc_gal_isom.
 DR InterPro; IPR005900; Phosphogluconac.
 DR Pfam; PF01182; Glucosamine_iso; 1.
 DR TIGRFAMS; TIGR01198; pgl; 1.
 KW Hydrolase; Complete proteome.
 SQ SEQUENCE 220 AA; 25525 MW; 9B0FD07E501E60C3 CRC64;
 Query Match 34.8%; Score 48; DB 1; Length 220;
 Best Local Similarity 34.8%; Pred. No. 5.5;

Search completed: September 15, 2003, 17:22:59
Job time : 7.12857 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 ; Search time 28.2286 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-1

Perfect score: 138
Sequence: 1 NLNAAQRYGRELRLRMSDFEGSFKGL 26

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SPTREMBL23:*

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mhc.*
- 8: sp_organelle.*
- 9: sp_phage.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*
- 15: sp_virus.*
- 16: sp_bacteriap.*
- 17: sp_archaeap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	87	63.0	146	13	Q919N2
2	53	38.4	196	16	Q8VJS3
3	53	38.4	223	16	Q10843
4	53	38.4	946	11	Q8K016
5	52	37.7	471	17	Q8ZY71
6	51	37.0	231	10	Q9SEGO
7	51	37.0	232	10	Q9SQ20
8	51	37.0	232	10	Q9SQ22
9	51	37.0	232	10	Q9SQ17
10	51	37.0	232	10	Q9SQ19
11	51	37.0	232	10	Q9SQ21
12	51	37.0	232	10	Q8LB79
13	51	37.0	232	10	Q8SQ16
14	51	37.0	232	10	Q9SQ15
15	51	37.0	232	10	Q9SQ18
16	51	37.0	232	10	Q9SQ18

17	50.5	36.6	904	2	Q9KGW3	Q9kgw3 pseudomonas
18	50.5	36.6	909	16	Q8EI34	Q8ei34 shewanella
19	50	36.2	168	11	Q8K316	Q8k316 mus musculus
20	50	36.2	283	15	Q37056	Q37056 chimpanzee
21	50	36.2	374	17	Q9H9Z9	Q9h9z9 halobacteri
22	50	36.2	516	10	Q9SSP5	Q9ssp5 arabidopsis
23	50	36.2	851	11	Q8K2H3	Q8k2h3 mus musculus
24	49.5	35.9	153	5	Q9UB33	Q9ub33 anopheles g
25	49.5	35.9	401	5	Q97407	Q97407 anopheles g
26	49.5	35.9	505	8	Q47148	Q47148 menziesia c
27	49.5	35.9	506	8	Q47149	Q47149 rhododendro
28	49.5	35.9	506	8	Q47171	Q47171 rhododendro
29	49.5	35.9	506	8	Q63960	Q63960 rhododendro
30	49.5	35.9	506	8	Q62982	Q62982 rhododendro
31	49.5	35.9	506	8	Q62975	Q62975 rhododendro
32	49.5	35.9	506	8	Q62972	Q62972 rhododendro
33	49.5	35.9	506	8	Q62989	Q62989 rhododendro
34	49.5	35.9	506	8	Q62978	Q62978 rhododendro
35	49.5	35.9	506	8	Q47155	Q47155 rhododendro
36	49.5	35.9	506	8	Q47152	Q47152 rhododendro
37	49.5	35.9	506	8	Q47173	Q47173 rhododendro
38	49.5	35.9	506	8	Q62990	Q62990 rhododendro
39	49.5	35.9	506	8	Q62974	Q62974 rhododendro
40	49.5	35.9	506	8	Q62993	Q62993 menziesia m
41	49.5	35.9	506	8	Q47170	Q47170 rhododendro
42	49.5	35.9	506	8	Q47174	Q47174 rhododendro
43	49.5	35.9	506	8	Q62983	Q62983 rhododendro
44	49.5	35.9	506	8	Q62980	Q62980 rhododendro
45	49.5	35.9	506	8	Q62981	Q62981 rhododendro

ALIGNMENTS

RESULT 1

Q919N2 PRELIMINARY; PRT; 146 AA.

AC Q919N2: 01-OCT-2000 (TREMREL. 15, Created)

DT 01-DEC-2001 (TREMREL. 19, Last sequence update)

DT 01-OCT-2002 (TREMREL. 22, Last annotation update)

DE Bad.

GN BAD.

OS Brachydanio rerio (Zebrafish) (Danio rerio).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;

OC Cyprinidae; Danio.

OX NCBI_TaxID=7955;

RN [1]

RP MEDLINE=20373792; PubMed=10917738;

RA Inohara N., Nunez G.;

RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish."

RL Cell Death Differ. 7:509-510(2000).

DR EMBL; AF231017; AAF66962.2; -

DR HSSP; Q92934; 165J

DR ZFIN; ZDB-GENE-000616-1; bad.

SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;

Query Match 63.0%; Score 87; DB 13; Length 146;
Best Local Similarity 65.2%; Pred. No. 5.3e-05;
Matches 15; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

Qy 2 LWAQRYGRELRLRMSDFEGSFK 24
||||:|||||:|

Db 89 LWAAKYQQLRLRMSDFEGKMK 111

RESULT 2

Q8VJS3 PRELIMINARY; PRT; 196 AA.

ID Q8VJS3

AC Q8VJS3;


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DT 01-MAR-2002 (TremBLrel. 20, Created)
DT 01-MAR-2002 (TremBLrel. 20, Last sequence update)
DT 01-JUN-2002 (TremBLrel. 21, Last annotation update)
DE IS1607, transposase.
GN MT2070.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W.;
RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."; to the EMBL/GenBank/DBJ databases.
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE007058; AAK46348.1; -.
DR TIGR; MT2070; -.
DR InterPro; IPR003346; Transposase_20.
DR Pfam; PF02371; Transposase_20; 1.
DR SEQUENCE 196 AA; 21349 MW; C145A8D836FD5C2D CRC64;
SQ
Query Match 38.4%; Score 53; DB 16; Length 196;
Best Local Similarity 58.8%; Pred. No. 7.7;
Matches 10; Conservative 1; Mismatches 6; Indels 0; Gaps 0;
QY 1 NLWAAORYGRELRLMSD 17
| | | | | | | | | | | | | | | | | | | | |
Db 134 NLWADRYNRAIARGHD 150
| | | | | | | | | | | | | | | | | | | | |

RESULT 3
Q10843
ID Q10843 PRELIMINARY; PRT; 223 AA.
AC Q10843;
DT 01-NOV-1998 (TremBLrel. 08, Created)
DT 01-NOV-1998 (TremBLrel. 08, Last sequence update)
DT 01-MAR-2002 (TremBLrel. 20, Last annotation update)
DE Hypothetical protein Rv2014.
GN Rv2014 OR MFCY39.03C.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies K., Devlin K., Feldwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
CC -!- SIMILARITY: TO M.PARATUBERCULOSIS IS900.
DR EMBL; Z74025; CAAB8415.1; -.
DR TuberculList; Rv2014; -.
DR InterPro; IPR003346; Transposase_20.
DR Pfam; PF02371; Transposase_20; 1.
DR Hypothetical protein; Complete proteome.
KW SEQUENCE 223 AA; 24132 MW; 70456750017FEF37 CRC64;
SQ
Query Match 38.4%; Score 53; DB 16; Length 223;
Best Local Similarity 58.8%; Pred. No. 8.9;

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Matches 10; Conservative 1; Mismatches 6; Indels 0; Gaps 0;
QY 1 NLWAAORYGRELRLMSD 17
| | | | | | | | | | | | | | | | | | | | |
Db 165 NLWADRYNRAIARGHD 181
| | | | | | | | | | | | | | | | | | | | |

RESULT 4
Q8X016
ID Q8X016 PRELIMINARY; PRT; 946 AA.
AC Q8X016;
DT 01-OCT-2002 (TremBLrel. 22, Created)
DT 01-OCT-2002 (TremBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TremBLrel. 23, Last annotation update)
DE Inter-alpha trypsin inhibitor, heavy chain 2.
GN ITIH2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Strausberg R.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC034341; AAH34341.1; -.
DR MGD; MGI:96619; Itih2.
DR InterPro; IPR006587; VIT.
DR InterPro; IPR002035; VWF_A.
DR SMART; SM00609; VIT; 1.
DR SMART; SM00327; VWA; 1.
DR PROSITE; PS50234; VWF_A; 1.
DR SEQUENCE 946 AA; 105945 MW; 8B17DBA71B85EC5C CRC64;
SQ
Query Match 38.4%; Score 53; DB 11; Length 946;
Best Local Similarity 34.6%; Pred. No. 44;
Matches 9; Conservative 5; Mismatches 12; Indels 0; Gaps 0;
QY 1 NLWAAORYGRELRLMSDSEFGSKGL 26
| : | : | : | : | : | : | : | : | : | : | : | : |
Db 212 NVWIMEPQGMRLHVPDFEGHFGGV 237
| : | : | : | : | : | : | : | : | : | : | : | : |

RESULT 5
Q8ZV71
ID Q8ZV71 PRELIMINARY; PRT; 471 AA.
AC Q8ZV71;
DT 01-MAR-2002 (TremBLrel. 20, Created)
DT 01-MAR-2002 (TremBLrel. 20, Last sequence update)
DT 01-MAR-2003 (TremBLrel. 23, Last annotation update)
DE Hypothetical protein PAE0922.
GN PAE0922.
OS Pyrobaculum aerophilum.
OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;
OC Thermoproteaceae; Pyrobaculum.
OX NCBI_TaxID=13773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IM2 / ATCC 51768 / DSM 7523;
RX PubMed=11792869;
RA Fitz-Gibbon S.T., Ladner H., Kim U.-J., Stetter K.O., Simon M.I.,
RA Miller J.H.;
RA "Genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum
RT aerophilum.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989(2002).
DR EMBL; AB009793; AAL63125.1; -.
DR InterPro; IPR006638; Elp3.
DR InterPro; IPR000182; GCN5acetyltransf.
DR Pfam; PF00583; Acetyltransf; 1.
DR SMART; SM00729; Elp3; 1.
DR Hypothetical protein; Complete proteome.
KW SEQUENCE 471 AA; 52952 MW; 3B1E36E8AE2EF0A CRC64;
SQ

```

Query Match 37.7%; Score 52; DB 17; Length 471;
Best Local Similarity 41.7%; Pred. No. 29;
Matches 10; Conservative 3; Mismatches 11; Indels 0; Gaps 0;

QY 3 WAAQRYGRLRRMSDFEGSKGL 26
DB 404 WOHSGNGRLMLAEIAGEFGL 427

RESULT 6
Q9SEGO PRELIMINARY; PRT; 231 AA.
AC Q9SEGO:
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-MAY-2000 (Tremblrel. 13, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE Apetala3 (Fragment).
OS Arabidopsis lyrata.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=59689;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=99404148; PubMed=10474900;
RA Lawton-Rauh A.L., Buckler E.S. IV, Purugganan M.D.;
RT "Patterns of molecular evolution among paralogous floral homeotic
genes.";
RL Mol. Biol. Evol. 16:1037-1045(1999).
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS
DR EMBL; AF143380; AAF25590.1; -.
DR HSSP; P11746; LMNM.
DR InterPro; IPR002487; TF_Kbox.
DR SMART; SM00432; MADS; 1.
DR Pfam; PF01486; K-box; 1.
DR PRINTS; PR00319; SRF-TF; 1.
DR PROSITE; PS00432; MADS_BOX_1.
DR PROSITE; PS00066; MADS_BOX_2; 1.
DR DNA-binding; Nuclear protein; Transcription; Transcription regulation.
KW Arabidopsis thaliana (Mouse-ear cross).
SQ SEQUENCE 231 AA; 27176 MW; A67CAE1EBD8F7A CRC64;

Query Match 37.0%; Score 51; DB 10; Length 231;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDFEGSKF 24
DB 107 QRLGCLDELDIQELRLDEMENTFK 133

RESULT 7
Q9SQ20 PRELIMINARY; PRT; 232 AA.
AC Q9SQ20:
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-MAY-2000 (Tremblrel. 13, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=cv. Corscalla;
RX MEDLINE=99126449; PubMed=9927474;
RA Purugganan M.D., Suddith J.I.;

RT "Molecular population genetics of floral homeotic loci. Departures
from the equilibrium-neutral model at the APETALA3 and PISTILLATA
genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS
DR EMBL; AF115806; AAD51895.1; -.
DR HSSP; P11746; LMNM.
DR InterPro; IPR002487; TF_Kbox.
DR SMART; SM00432; MADS_BOX_1; 1.
DR PROSITE; PS00350; MADS_BOX_2; 1.
DR PROSITE; PS00066; MADS_BOX_2; 1.
DR DNA-binding; Nuclear protein; Transcription; Transcription regulation.
KW Arabidopsis thaliana (Mouse-ear cross).
SQ SEQUENCE 232 AA; 27342 MW; BDFDC59B73F4601 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDFEGSKF 24
DB 107 QRLGCLDELDIQELRLDEMENTFK 133

RESULT 8
Q9SQ22 PRELIMINARY; PRT; 232 AA.
AC Q9SQ22:
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-MAY-2000 (Tremblrel. 13, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=cv. Li-8;
RX MEDLINE=99126449; PubMed=9927474;
RA Purugganan M.D., Suddith J.I.;

RT "Molecular population genetics of floral homeotic loci. Departures
from the equilibrium-neutral model at the APETALA3 and PISTILLATA
genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS
DR EMBL; AF115801; AAD51890.1; -.
DR HSSP; P11746; LMNM.
DR InterPro; IPR002487; TF_Kbox.
DR SMART; SM00432; MADS_BOX_1; 1.
DR PROSITE; PS00350; MADS_BOX_2; 1.
DR PROSITE; PS00066; MADS_BOX_2; 1.
DR DNA-binding; Nuclear protein; Transcription; Transcription regulation.
KW Arabidopsis thaliana (Mouse-ear cross).
SQ SEQUENCE 232 AA; 27267 MW; 42A852D697E22A65 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDFEGSKF 24


```

RESULT 12
Q8LB79 ID Q8LB79 PRELIMINARY; PRT; 232 AA.
AC EMBL; AF115810; AAD51899.1; -.
DT 01-OCT-2002 (TRENBLrel. 22, Created)
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Floral homeotic protein APETALA3 (AP3).
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA Haas B.J., Volfovsky N., Town C.D., Troupkhan M., Alexandrov N.,
RA Feldmann K.A., Flavell R.B., White O., Salzberg S.L.;
RT "Full-length messenger RNA sequences greatly improve genome
RT annotation.";
RL Genome Biol. 0:0-0(2002).
RN [2]
RP SEQUENCE FROM N.A.
RA Brover V., Troupkhan M., Alexandrov N., Lu Y.-P., Flavell R.,
RA Feldmann K.;
RT "Full-length cDNA from Arabidopsis thaliana.";
RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AY087369; AAM64919.1; -.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27339 MW; CC90703F959CFAD5 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEFGSKF 24
DB 107 QRLGCLDKLDIQELRLLEDEMENTFK 133

RESULT 13
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AC Q9SQ16;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=cv. Lisse;
RA MEDLINE=99126449; PubMed=9927474;
RA Purganan M.D., Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
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DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27300 MW; 5CA05FDA4F824DF0 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEFGSKF 24
DB 107 QRVGCLDKLDIQELRLLEDEMENTFK 133

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RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
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DR HSP; P11746; IMNM.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27314 MW; DB8CALFC835557D6 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEFGSKF 24
DB 107 QRLGCLDKLDIQELRLLEDEMENTFK 133

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AC Q9SQ15;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=cv. Kas-1;
RA MEDLINE=99126449; PubMed=9927474;
RA Purganan M.D., Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AF115812; AAD51901.1; -.
DR HSP; P11746; IMNM.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27300 MW; 5CA05FDA4F824DF0 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEFGSKF 24
DB 107 QRVGCLDKLDIQELRLLEDEMENTFK 133

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RESULT 15
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AC Q9S7Q3;
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DT 01-MAY-2000 (TREMELrel. 13, Last sequence update)
DT 01-OCT-2002 (TREMELrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsiis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. GR-3, and cv. CHI-1;
RX MEDLINE=99126449; PubMed=9927474;
RA Purganan M.D., Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
RT genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AF115803; AAD51892.1; -.
DR EMBL; AF115798; AAD51887.1; -.
DR HSP; P11746; 1MNM.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRF-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27340 MW; 6690703F9F9CFD63 CRC64;

Query Match 37.0%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 18;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRRMSDEFEQSPK 24
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Db 107 QRLGECLEKLDIQELRLREDEMENTFK 133

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Search completed: September 15, 2003, 17:25:45
Job time : 30.2286 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 23, 2003, 09:43:16 ; Search time 86 Seconds
(without alignments)
49.833 Million cell updates/sec

Title: US-09-544-664B-2

Perfect score: 142

Sequence: 1 NLWRAQFVGRLEKMSDFVDFKKGIL (27)

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	DB ID	Description
1	137	96.5	166	BBC6 protein for r
2	137	96.5	168	Human Bcl-xL/Bcl-2
3	137	96.5	168	Human cell prolif
4	137	96.5	168	Human BAD mutant a
5	137	96.5	168	Human BAD protein.
6	137	96.5	168	Amino acid sequenc
7	137	96.5	168	Human BAD protein
8	137	96.5	201	Human ovarian anti
9	127	89.4	25	PTPC-interacting T

10	127	89.4	25	23	ABG78484	Mutant Bcl2 compet
11	127	89.4	25	23	AAU78610	Human Bad peptide
12	124	87.3	25	23	ABG78490	Mutant Bcl2 compet
13	124	87.3	25	23	AAU78617	Human Bad peptide
14	123	86.6	25	23	ABG78488	Mutant Bcl2 compet
15	123	86.6	25	23	ABG78489	Mutant Bcl2 compet
16	123	86.6	25	23	AAU78615	Human Bad peptide
17	123	86.6	25	23	AAU78616	Human Bad peptide
18	122	85.9	24	23	AAU78627	Human Bad peptide
19	122	85.9	25	23	ABG78486	Mutant Bcl2 compet
20	122	85.9	25	23	ABG78493	Mutant Bcl2 compet
21	122	85.9	25	23	ABG78497	Mutant Bcl2 compet
22	122	85.9	25	23	AAU78612	Human Bad peptide
23	122	85.9	25	23	AAU78620	Human Bad peptide
24	122	85.9	25	23	AAU78624	Human Bad peptide
25	121	85.2	25	23	ABG78485	Mutant Bcl2 compet
26	121	85.2	25	23	ABG78492	Mutant Bcl2 compet
27	121	85.2	25	23	AAU78611	Human Bad peptide
28	119	83.8	25	23	AAU78619	Human Bad peptide
29	119	83.8	25	23	ABG78491	Mutant Bcl2 compet
30	119	83.8	25	23	ABG78498	Mutant Bcl2 compet
31	119	83.8	25	23	AAU78618	Human Bad peptide
32	119	83.8	25	23	AAU78625	Human Bad peptide
33	117	82.4	23	23	AAU78628	Human Bad peptide
34	116	81.7	26	21	AAV96321	Mammalian Bad Bcl-
35	116	81.7	26	22	ABG70371	BAD BH3 consensus
36	113	79.6	25	23	ABG78487	Mutant Bcl2 compet
37	113	79.6	25	23	AAU78613	Human Bad peptide
38	112	78.9	25	23	ABG78495	Mutant Bcl2 compet
39	112	78.9	25	23	AAU78622	Human Bad peptide
40	111	78.2	22	23	AAU78629	Human Bad peptide
41	110	77.5	25	23	ABG78496	Mutant Bcl2 compet
42	110	77.5	25	23	AAU78623	Human Bad peptide
43	109	76.8	26	21	AAAB37001	Bcl2 polypeptide B
44	109	76.8	26	21	AAAB37002	Bcl2 polypeptide B
45	109	76.8	27	21	AAAB37003	Bcl2 polypeptide B

ALIGNMENTS

RESULT 1

AAW32476

ID AAW32476 standard; Protein; 166 AA.

XX AAW32476;

AC AAW32476;

DT 15-JAN-1998 (first entry)

DE BBC6 protein for regulating cell death.

KW BBC6 gene; cell death; cell cycle; Bcl2; human.

XX Homo sapiens.

XX US5663316-A.

XX 02-SEP-1997.

XX 18-JUN-1996; 96US-0665617.

XX 18-JUN-1996; 96US-0665617.

XX (CLON-) CLONTECH LAB INC.

XX Xudong Y;

XX WPI: 1997-447980/41.

XX N-PSDB; AAT91561.

XX Isolated BBC6 gene - encodes a protein that regulates cell death

XX through interaction with Bcl-2

PS Claim 1; Column 11-12; 7pp; English.

XX The present sequence represents a protein of 166 amino acids. The
CC sequence is disclosed as being a protein called Bcl-2 which regulates
CC cell death through interaction with Bcl-2. The DNA may be used for the
CC production of the recombinant protein, which can be used in unspecified
CC therapeutic or diagnostic procedures, as a molecular weight marker, and
CC to raise antibodies that can be used in unspecified diagnostic or
CC therapeutic applications and to reduce or eliminate the biological
CC activity of the Bcl-2 protein in vivo.

XX Sequence 166 AA;

Query Match 96.5%; Score 137; DB-18; Length 166;
Best Local Similarity 96.3%; Pred. No. 1-1e-12;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFVDSFKKGL 27
DB 101 NLWAAQRYGRELRLMSDEFVDSFKKGL 127

RESULT 2

AAW55779
ID AAW55779 standard; Protein; 168 AA.

XX AAW55779;

XX 17-JUL-1998 (first entry)

XX Human Bcl-xL/Bcl-2 associated death promoting polypeptide.

XX Human; Bcl-xL/Bcl-2 associated death promoting polypeptide; Bad;
KW programmed cell death; apoptosis.

XX Homo sapiens.

XX WO9812328-A2.

XX 26-MAR-1998.

XX 18-SEP-1997; 97WO-US16991.

XX 20-SEP-1996; 96US-0717123.

XX (IDUN-) IDUN PHARM INC.

XX Horne WA, Oltersdorf T;

XX WPI; 1998-21267/19.

XX N-PSDB; AAV25877.

XX Bad gene mediating apoptosis - used to develop products for treating
XX e.g. neurodegenerative disease, cancers or autoimmune disease

PS Claim 8; Fig 1; 41pp; English.

XX The present sequence is the human Bcl-xL/Bcl-2 associated
CC death promoting polypeptide, Bad, the binding of which to Bcl-xL
CC results in the induction of programmed cell death, i.e. apoptosis.
CC Bad can be used in screening assays for compounds to treat or
CC prevent diseases characterised by apoptotic cell death, such as
CC neurodegenerative disorders, e.g. Alzheimer's and Parkinson's
CC disease, amyotrophic lateral sclerosis, retinitis pigmentosa and
CC cerebellar degeneration, and myelodysplastic syndromes, e.g.

XX aplastic anaemia and ischaemic injury including myocardial
CC infarction, stroke and reperfusion injury. Assays can also be
CC used to obtain apoptosis enhancing compounds to treat or prevent
CC diseases characterised by the loss of apoptotic cell death, such as
CC cancers, e.g. lymphoma and hormone dependent tumours, autoimmune
CC diseases, e.g. systemic lupus erythematosus and immune-mediated
CC glomerulonephritis and viral infections, e.g. herpesvirus,
XX poxvirus or adenovirus infection. Bad can also be used for

CC detection and diagnosis.

XX Sequence 168 AA;

Query Match 96.5%; Score 137; DB-18; Length 168;
Best Local Similarity 96.3%; Pred. No. 1-1e-12;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFVDSFKKGL 27
DB 103 NLWAAQRYGRELRLMSDEFVDSFKKGL 129

RESULT 3

AA13512
ID AA13512 standard; protein; 168 AA.

XX AA13512;

XX 02-NOV-2000 (first entry)

XX Human cell proliferation protein APOP-1.

XX Human; cell proliferation; APOP-1; cancer; inflammation; infection;
KW trauma; neurodegenerative disease; ischaemic injury; wasting disease.

XX Homo sapiens.

XX US6080847-A.

XX 27-JUN-2000.

XX 04-DEC-1997; 97US-0985335.

XX 04-DEC-1997; 97US-0985335.

XX (INCY-) INCYTE PHARM INC.

XX Corley NC, Hillman JL, Yue H, Lal P, Shah P;

XX WPI; 2000-451230/39.

XX N-PSDB; AAA63332.

XX Novel polynucleotide and polypeptide sequences of proteins associated
XX with cell proliferation for diagnosis, prevention and treatment of e.g.
XX cancer, acquired immunodeficiency syndrome, and Parkinson's disease -

PS Example 8; Fig 1; 58pp; English.

XX The present sequence is the human APOP-1 protein. This protein, which
CC shares structural and chemical homology with Bcl-2, is involved in cell
CC proliferation. Its coding sequence was isolated by screening a synovial
CC tissue cDNA library using a computer search for amino acid sequence
CC alignments. The gene and protein can be used in the treatment of various
CC cancers, disorders with associated inflammation such as Addison's
CC disease, adult respiratory distress syndrome, allergies, anaemia, asthma,
CC atherosclerosis, Crohn's disease, ulcerative colitis, diabetes mellitus,
CC emphysema, glomerulonephritis, gout, Graves' disease, irritable bowel
CC syndrome, lupus erythematosus, multiple sclerosis, myasthenia gravis,
CC myocardial or pericardial inflammation, osteoporosis, rheumatoid
CC arthritis, Sjogren's syndrome and autoimmune thyroiditis, complications
CC of cancer, haemodialysis and extracorporeal circulation, infections,
CC trauma, disorders with associated apoptosis including AIDS and other
CC infectious and genetic immunodeficiencies, neurodegenerative diseases
CC such as Alzheimer's disease and Parkinson's disease, ischaemic injuries
CC such as myocardial infarction, and wasting diseases including cachexia.

XX Sequence 168 AA;

Query Match 96.5%; Score 137; DB-18; Length 168;
Best Local Similarity 96.3%; Pred. No. 1-1e-12;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFVDSFKKGL 27
 DB 103 NLWAAQRYGRELRRMSDEFVDSFKKGL 129

RESULT 4

AAB70368
 ID AAB70368 standard; protein; 168 AA.

XX AC AAB70368;

XX DT 02-MAY-2001 (first entry)

XX DE Human BAD mutant amino acid sequence SEQ ID NO:1.

XX KW Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 KW immunostimulant; neuroprotective; nootropic; antiischaemic; vulnerary;
 KW cytosolic; antiviral; antiarthritic; antiinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.

XX OS Homo sapiens.
 XX SY Synthetic.

PN W0200110888-A1.

XX PD 15-FEB-2001.

XX PF 30-MAY-2000; 2000WO-US11864.

XX PR 28-MAY-1999; 99US-0136783.

XX PA (APOP-) APOPTOSIS TECHNOLOGY INC.

XX PI Zhou X;

XX DR WPI; 2001-138734/14.

XX PT New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -

XX PS Claim 1; Page 147; 157pp; English.

XX CC The present invention describes an isolated or synthetic polypeptide
 CC (I) comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
 CC nootropic, antiischaemic, vulnerary, cytosolic, antiviral,
 CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed human BAD mutant amino acid sequence from the present invention.

XX SQ Sequence 168 AA;

Query Match 96.5%; Score 137; DB 22; Length 168;
 Best Local Similarity 96.3%; Pred. No. 1.1e-12;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFVDSFKKGL 27

DB 103 NLWAAQRYGRELRRMSDEFVDSFKKGL 129

RESULT 5

AAB48287
 ID AAB48287 standard; protein; 168 AA.

XX AC AAB48287;

XX DT 02-APR-2001 (first entry)

XX DE Human Bad protein.

XX KW S-phase kinase associated protein; SKP1; SKP2; SKP2-like protein; 2F;
 KW CUL-1; cullin; CDC53; p27; cyclin E; Max; Mad; c-Myc; MDM2; p53; Bax;
 KW Bad; Bcl-2; tumour; cytosolic.

XX OS Homo sapiens.

XX PN W0200075184-A1.

XX PD 14-DEC-2000.

XX PF 05-JUN-2000; 2000WO-US15449.

XX PR 04-JUN-1999; 99US-0137494.

XX PA (UYTA) UNIV YALE.

XX PI Zhang H, Tsvetkov LM, Kondo T;

XX DR WPI; 2001-061703/07.

XX DR N-ESDB; AAC84599.

XX PT Modulating polypeptide levels in a cell, diagnosing and treating tumor,
 PT involves altering levels of proteins such as S-phase kinase associated
 PT proteins 1, 2 and cullin/CDC53 proteins -

XX PS Claim 5; Page 102-103; 162pp; English.

XX CC The invention relates to methods of altering the polypeptide levels in a
 CC cell, using proteins selected from S-phase kinase associated proteins 1
 CC and 2 (SKP1, SKP2), SKP2-like proteins (2F) and CUL-1 (a member of the
 CC cullin/CDC53 family of proteins). The method is useful for altering the
 CC level of p27, cyclin E, Max, Mad, c-Myc, MDM2, p53, Bax, Bad or Bcl-2
 CC polypeptide in a cell. SKP2 and SKP2-like protein levels are useful for
 CC detecting tumours, and in monitoring tumor treatment in a mammal. Agents
 CC that modulate interactions between SKP and target proteins are useful for
 CC treating tumours.

XX SQ Sequence 168 AA;

Query Match 96.5%; Score 137; DB 22; Length 168;
 Best Local Similarity 96.3%; Pred. No. 1.1e-12;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFVDSFKKGL 27

DB 103 NLWAAQRYGRELRRMSDEFVDSFKKGL 129

RESULT 6

AAG67688
 ID AAG67688 standard; Protein; 168 AA.

XX AC AAG67688;

XX DT 26-NOV-2001 (first entry)

XX DE Amino acid sequence of protein associated with cell proliferation-1.

XX KW Human; cell proliferation; APOP-1; APOP-2; APOP-3; apoptosis; cancer;

KW brain cancer; breast cancer; Alzheimer's disease; Parkinson's disease;
 KW inflammation; allergy; gout; osteoarthritis; bronchitis.
 XX Homo sapiens.
 XX Key Location/Qualifiers
 FT Modified-site 10..13
 FT /note= "potential casein kinase II phosphorylation site"
 FT Modified-site 16..19
 FT /note= "potential casein kinase II phosphorylation site"
 FT Modified-site 34..36
 FT /note= "potential protein kinase C phosphorylation site"
 FT Modified-site 80..83
 FT /note= "potential casein kinase II phosphorylation site"
 FT Modified-site 115..118
 FT /note= "potential CAMP- and CGMP-dependent protein
 FT kinase phosphorylation site"
 FT Modified-site 124..126
 FT /note= "potential protein kinase C phosphorylation site"
 FT Modified-site 153..156
 FT /note= "potential casein kinase II phosphorylation site"
 XX US6281334-B1.
 XX 28-AUG-2001.
 XX 30-SEP-1999; 99US-0410372.
 XX 04-DEC-1997; 97US-0985335.
 XX (INCY-) INCYTE GENOMICS INC.....
 XX Hallman JL, Yue H, Lal P, Shah P, Corley NC;
 DR WPI: 2001-569961/64.
 DR N-PSDB; AAH78430.
 XX New polypeptides associated with cell proliferation, useful for
 FT preventing or treating cancer (e.g. brain cancer), a disorder
 FT associated with an increase in apoptosis (e.g. Alzheimer's disease) or
 FT inflammation (e.g. gout) -
 XX Example; Fig 1A-C; 59pp; English.
 PS The present sequence represents a human protein which is associated
 CC with cell proliferation, designated APOP-1. The specification also
 CC describes APOP-2 and APOP-3. The APOP polypeptides are useful for
 CC diagnosing, preventing or treating disorders associated with abnormal
 CC cell proliferation and apoptosis. The polypeptides and composition are
 CC particularly useful for treating or preventing cancer (e.g. brain or
 CC breast cancer), a disorder associated with an increase in apoptosis
 CC (e.g. Alzheimer's disease or Parkinson's disease) or inflammation
 CC (e.g. allergies, gout, osteoarthritis or bronchitis).
 XX Sequence 168 AA;
 SQ Query Match 96.5%; Score 137; DB 22; Length 168;
 Best Local Similarity 96.3%; Pred. No. 1,1e-12;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDEFVDSFKGL 27
 DB 103 NLWAAQRYGRELRRMSDEFVDSFKGL 129
 RESULT 7
 ABR39081
 ID ABR39081 standard; Protein; 168 AA.
 XX ABR39081;
 AC ABR39081;
 XX 10-MAY-2003 (first entry)
 DT
 XX

DE Human BAD protein SEQ ID NO:2.
 XX Human; BAD; herpes simplex virus; HSV; US3; herpes virus; apoptosis;
 KW virucide; infection.
 XX Homo sapiens.
 OS WO2003012049-A2.
 PN 13-FEB-2003.
 PD 31-JUL-2002; 2002WO-US24177.
 XX 31-JUL-2001; 2001US-308929P.
 PF (UYCH-) UNIV CHICAGO.
 PR Munger J, Roizman B;
 PA WPI: 2003-248168/24.
 FT N-PSDB; ABZ81200.
 DR Inducing apoptosis in a cell infected with herpes simplex virus, HSV,
 XX by administering to the cell, a composition comprising an agent that
 XX inhibits phosphorylation of pro-apoptotic polypeptide BAD by HSV US3
 PS Claim 15; Page 166-167; 192pp; English.
 XX The present invention describes a method (M1) for inducing apoptosis in
 CC a cell infected with herpes simplex virus (HSV), which comprises
 CC administering to the cell, a composition having an agent that inhibits
 CC phosphorylation of pro-apoptotic polypeptide BAD by HSV US3. Also
 CC described is a method (M2) for treating a patient infected with HSV, by
 CC administering to the patient, a composition comprising a peptide
 CC comprising a sequence of 4-100 continuous amino acids of a 168 residue
 CC amino acid sequence (see ABR39081), where the peptide comprises ser112,
 CC ser135, or ser155, or their combinations. BAD has virucide activity.
 CC M1 is useful for inducing apoptosis in a cell infected with HSV, where
 CC the cell is in a human. M2 is useful for treating a patient infected
 CC with HSV. The present sequence represents human BAD, which is used in
 CC the exemplification of the present invention.
 XX Sequence 168 AA;
 SQ Query Match 96.5%; Score 137; DB 24; Length 168;
 Best Local Similarity 96.3%; Pred. No. 1,1e-12;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRRMSDEFVDSFKGL 27
 DB 103 NLWAAQRYGRELRRMSDEFVDSFKGL 129
 RESULT 8
 ABP41630
 ID ABP41630 standard; Protein; 201 AA.
 XX AC ABP41630;
 XX 22-AUG-2002 (first entry)
 DT
 XX Human ovarian antigen HCE4K28, SEQ ID NO:2762.
 DE Human; ovarian antigen; ovary; ovarian; breast; cancer; tumour;
 KW ovarian cancer; breast cancer; tumour; reproductive system disorder;
 KW infertility; pregnancy disorder; anovulation; polycystic ovary syndrome;
 KW PCOS; ovarian cyst; dysmenorrhea; endocrine disorder; infection;
 KW inflammatory condition; immune disorder; blood disorder;
 KW cardiovascular disorder; respiratory disorder; neurological disorder;
 KW gastrointestinal disorder; urinary system disorder; drug screening;
 KW gene therapy; chromosome mapping; forensic analysis;
 KW antibody preparation; cytostatic; immunomodulatory; neuroprotective;
 KW antiinflammatory; gynaecological; reproductive.

XX OS Homo sapiens.
 XX FN W0200200677-A1.
 XX PD 03-JAN-2002.
 XX PF 07-JUN-2001; 2001WO-US18569.
 XX PR 07-JUN-2000; 2000US-209467P.
 XX PA (HUMA-) HUMAN GENOME SCI INC.
 XX PI Birse CE, Rosen CA;
 XX DR WPI; 2002-147878/19.
 XX DR N-PSDB; ABQ54707.
 XX PT Isolated nucleic acid molecules encoding novel ovarian polypeptides,
 XX PT useful in the prevention, treatment and diagnosis of cancer (e.g.
 XX PT ovarian cancer), immune disorders, cardiovascular disorders and
 XX PT neurological diseases -
 XX PS Claim 11; SEQ ID No 2762; 2922pp; English.
 XX CC The invention relates to 2175 novel human ovarian antigens (ABP41054-
 CC ABP43228) and to cDNAs encoding them (ABQ54131-ASQ56305), and also
 CC encompasses polypeptides 90% identical and polynucleotides 95% identical
 CC to the sequences of the invention. The invention additionally relates to
 CC recombinant vectors and host cells comprising human ovarian antigen
 CC polynucleotides, antibodies against human ovarian antigens, and the use
 CC of ovarian antigen polynucleotides and polypeptides in diagnosing,
 CC treating, prognosing or preventing various ovary and/or breast-related
 CC disorders. Such conditions include ovarian cancer and breast cancer, and
 CC metastatic tumours of ovarian or breast origin, reproductive system
 CC disorders (e.g., infertility, disorders of pregnancy, anovulation,
 CC polycystic ovary syndrome, ovarian cysts, and dysmenorrhoea), endocrine
 CC disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and toxic
 CC shock syndrome), inflammatory conditions (e.g., mastitis, oophoritis and
 CC vaginitis), immune disorders (e.g., congenital and acquired
 CC immunodeficiencies, autoimmune oophoritis, systemic lupus erythematosus),
 CC blood-related disorders (e.g., anaemia), cardiovascular disorders,
 CC respiratory disorders, neurological disorders, gastrointestinal disorders
 CC and urinary system disorders. Ovarian antigen polypeptides and
 CC polynucleotides may also be used in screening for compounds which
 CC modulate ovarian antigen expression or activity. The polynucleotides may
 CC further be used for gene therapy, chromosome mapping, in the
 CC identification of individuals and in forensic analysis, and the
 CC polypeptides may be used as food additives or to prepare antibodies
 CC useful in disease diagnosis, drug targeting and phenotyping. The present
 CC sequence represents a human ovarian antigen of the invention.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX SQ Sequence 201 AA;
 Query Match 96.58; Score 137; DB 23; Length 201;
 Best Local Similarity 96.3%; Pred. No. 1.3e-12;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 NLWAAQEGRELRMSDFVDSFKGL 27
 DB 111 NLWAAQEGRELRMSDFVDSFKGL 137
 RESULT 9
 ABP56161
 ID ABP56161 standard; peptide; 25 AA.
 XX AC ABP56161;
 XX DT 28-MAR-2003 (first entry)

XX DE PTPC-interacting TOX peptide #27.
 XX KW Mitochondrial membrane permeabilisation; mitochondrion; PTPC;
 KW permeability transition pore complex; virucide; neuroprotective;
 KW vasotropic; cytostatic; infection; cell death regulation; apoptosis;
 KW mitochondrial permeability transition pore complex modulator; cancer;
 KW apoptogenic; ischaemia; neurodegenerative disease; fulminant hepatitis.
 XX OS Synthetic.
 XX PN W0200261105-A2.
 XX PD 08-AUG-2002.
 XX PF 01-FEB-2002; 2002WO-EP01633.
 XX PR 02-FEB-2001; 2001US-265594P.
 XX PA (INSP INST PASTEUR
 XX PA (CNRS CENT NAT RECH SCI
 XX PI Edelman L, Jacotot E, Briand J;
 XX DR WPI; 2002-619260/66.
 XX PT New chimeric bifunctional molecules that target specific cells and
 PT regulate the apoptosis function of the permeability transition pore
 PT complex of the mitochondria, useful for treating or preventing e.g.
 PT cancer or ischemia -
 XX PS Claim 9; Page 11; 76pp; English.
 XX CC The present invention describes a chimeric bifunctional molecule (I)
 CC comprising at least a first functional molecule covalently linked to a
 CC second functional molecule, which is able to modulate the activity of
 CC the permeability transition pore complex (PTPC) of the mitochondria.
 CC (I) has the function of specifically targeting and entering a tissue
 CC cell population. The second functional molecule has the function of
 CC specifically targeting, and inducing or preventing the death of the
 CC cells by apoptosis by regulating the opening or the closing of the PTPC
 CC of the mitochondria or its fragment. (I) has virucide, neuroprotective,
 CC vasotropic and cytostatic activities, and can be used as a mitochondrial
 CC permeability transition pore complex (PTPC) modulator. (I) is useful for
 CC treating or preventing a pathological infection or disease. (I) is also
 CC useful for regulating cell death regulatory molecules, specifically the
 CC apoptogenic function of the PTPC, for treating e.g. cancer, ischaemia,
 CC neurodegenerative diseases, fulminant hepatitis or viral infections.
 CC The present sequence represents a PTPC-interacting TOX peptide which is
 CC given in the exemplification of the present invention.
 XX SQ Sequence 25 AA;
 Query Match 89.48; Score 127; DB 23; Length 25;
 Best Local Similarity 96.0%; Pred. No. 4.2e-12;
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 NLWAAQEGRELRMSDFVDSFKK 25
 DB 1 NLWAAQEGRELRMSDFVDSFKK 25
 RESULT 10
 ABG78484
 ID ABG78484 standard; Peptide; 25 AA.
 XX AC ABG78484;
 XX DT 15-NOV-2002 (first entry)
 XX DE Mutant Bcl2 competitive binding assay peptide #1.
 XX KW Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; mutelin.

CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
CC shown in the specification. The invention also comprises an assay for
CC identifying substances that bind to the Bcl-2 protein. The protein
CC sequences of the invention are useful in biological assays to identify
CC substances that block the ability of Bcl-2 to inhibit programmed cell
CC death or apoptosis. The present sequence represents a human Bcl2
CC peptide of the invention.
XX
SQ Sequence 25 AA;

Query Match 87.3%; Score 124; DB 23; Length 25;
Best Local Similarity 92.0%; Pred. No. 1.2e-11;
Matches 23; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFVDSFKK 25
||||| |||||||||
Db 1 NLWAAQRYGRELRRMSDFVDAFKK 25

RESULT 13
AAU78617

ID AAU78617 standard; Peptide; 25 AA.

XX AC AAU78617;

XX XX 18-JUN-2002 (first entry)

XX DE Human Bad peptide #17 which binds to a member of the Bcl-2 family.

XX KW Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
XX KW ischemic injury; suppressor; BH3 domain.

XX OS Homo sapiens.

XX EN W0200220568-A2.

XX PD 14-MAR-2002.

XX PF 04-SEP-2001; 2001WO-US7410.

XX PR 06-SEP-2000; 2000US-0656399.

XX PA (ABBO) ABBOTT LAB.

XX PI Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
XX PI Nettesheim DG, Swift KM, Matayoshi E, Zhang H;

XX DR WPI; 2002-292254/33.

XX PT New derivatives of Bad peptide, useful for identifying compounds that
XX PT bind to Bcl-2 proteins, potential agents for treating cancer and
XX PT degenerative disease .
XX PS Example 1; Page 14; 31pp; English.

XX CC The present invention relates to new peptides that are derived from a
XX CC wild-type human Bad peptide and are able to bind to a member of the
XX CC Bcl-2 protein family. The peptides are useful, when labelled, in
XX CC competitive/displacement assays for identifying substances that bind to
XX CC members of the Bcl-2 family and may induce or suppress apoptosis so are
XX CC potentially useful for treating cancer (inducers) or degenerative
XX CC diseases or ischemic injury (suppressors). The peptides of the invention
XX CC have high helix propensity, maintain the contacts of the wild-type Bad
XX CC peptide and, compared with the Bad peptide, may have better physical
XX CC properties, particularly solubility. The present sequence represents one
XX CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
XX CC from the BH3 domain of the human wild-type Bad peptide.
XX SQ Sequence 25 AA;

Query Match 87.3%; Score 124; DB 23; Length 25;
Best Local Similarity 92.0%; Pred. No. 1.2e-11;
Matches 23; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFVDSFKK 25
||||| |||||||||
Db 1 NLWAAQRYGRELRRMSDFVDAFKK 25

RESULT 14
ABG78488

ID ABG78488 standard; Peptide; 25 AA.

XX AC ABG78488;

XX XX 15-NOV-2002 (first entry)

XX DE Mutant Bcl2 competitive binding assay peptide #5.

XX KW Human; Bcl2; BclX1; programmed cell death; apoptosis; mutant; mitein.

XX OS Homo sapiens.

XX OS Synthetic.

XX PN W0200240530-A2.

XX PD 23-MAY-2002.

XX PF 15-NOV-2001; 2001WO-US45693.

XX PR 20-NOV-2000; 2000US-0716395.

XX PA (ABBO) ABBOTT LAB.

XX PI Fesik SW, Petros AM, Yoon H, Nettesheim DG;

XX DR WPI; 2002-490141/52.

XX PT New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
XX PT useful in biological assays to identify substances that block the
XX PT ability of Bcl-2 to inhibit programmed cell death or apoptosis .
XX PS Example 2; Page 17; 36pp; English.

XX CC This invention relates to a novel mutant protein which is derived from
XX CC a wild type human Bcl-2 protein. The mutant is created by replacing a
XX CC sequence of amino acid residues comprising a flexible loop from the wild
XX CC type Bcl-2 protein with an amino acid sequence comprising at least two
XX CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
XX CC shown in the specification. The invention also comprises an assay for
XX CC identifying substances that bind to the Bcl-2 protein. The protein
XX CC sequences of the invention are useful in biological assays to identify
XX CC substances that block the ability of Bcl-2 to inhibit programmed cell
XX CC death or apoptosis. The present sequence represents a human Bcl2
XX CC peptide of the invention.
XX SQ Sequence 25 AA;

Query Match 86.6%; Score 123; DB 23; Length 25;
Best Local Similarity 92.0%; Pred. No. 1.7e-11;
Matches 23; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFVDSFKK 25
||||| |||||||||
Db 1 NLWAAQRYGRELRRMSDFVDSFKK 25

RESULT 15
ABG78489

ID ABG78489 standard; Peptide; 25 AA.

XX AC ABG78489;

XX XX 15-NOV-2002 (first entry)

XX DE Mutant Bcl2 competitive binding assay peptide #6.

```
XX Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; mutain.
KW Homo sapiens.
OS Synthetic.
XX WO200240530-A2.
XX 23-MAY-2002.
XX 15-NOV-2001; 2001WO-US45693.
XX 20-NOV-2000; 2000US-0716395.
XX (ABBO ) ABBOTT LAB.
XX Pesik SW, Petros AM, Yoon H, Nettesheim DG;
XX WPI; 2002-490141/52.
XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
PT useful in biological assays to identify substances that block the
PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -
XX
XX Example 2; Page 17; 36pp; English.
XX This invention relates to a novel mutant protein which is derived from
CC a wild type human Bcl-2 protein. The mutant is created by replacing a
CC sequence of amino acid residues comprising a flexible loop from the wild
CC type Bcl-2 protein with an amino acid sequence comprising at least two
CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
CC shown in the specification. The invention also comprises an assay for
CC identifying substances that bind to the Bcl-2 protein. The protein
CC sequences of the invention are useful in biological assays to identify
CC substances that block the ability of Bcl-2 to inhibit programmed cell
CC death or apoptosis. The present sequence represents a human Bcl2
XX peptide of the invention.
XX
XX Sequence 25 AA;
XX
Query Match 86.6%; Score 123; DB 23; Length 25;
Best Local Similarity 92.0%; Pred. No. 1.7e-11;
Matches 23; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 NLWAGQRYGRELRLRMSDFVDSFKK 25
Search completed: September 23, 2003, 09:47:13
Job time : 87 secs
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OM protein - protein search, using sw model

Run on: September 23, 2003, 09:43:16 ; Search time 30 Seconds
(without alignments)
38.080 Million cell updates/sec

Title: US-09-544-664B-2
Perfect score: 142
Sequence: 1 NLWAAQYGRGLRMSDFVDFKGL 27

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents_AA:*
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2: /cgn2_6/ptodata/1/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/1/iaa/6A_COMB.pep.*
4: /cgn2_6/ptodata/1/iaa/6B_COMB.pep.*
5: /cgn2_6/ptodata/1/iaa/PCTUS_COMB.pep.*
6: /cgn2_6/ptodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	137	96.5	166	1	US-08-665-617-2
2	137	96.5	168	2	Sequence 2, Appli
3	137	96.5	168	3	Sequence 1, Appli
4	137	96.5	168	3	Sequence 7, Appli
5	137	96.5	168	3	Sequence 1, Appli
6	137	96.5	168	3	Sequence 2, Appli
7	137	96.5	168	4	US-09-375-257-2
8	109	76.8	204	1	US-08-333-565-2
9	109	76.8	204	2	US-08-661-479-2
10	109	76.8	204	2	US-08-733-505A-1
11	109	76.8	204	2	US-08-733-505A-12
12	109	76.8	204	2	US-08-733-505A-13
13	109	76.8	204	2	US-08-733-505A-14
14	106	74.6	204	2	US-08-717-123-3
15	106	74.6	204	4	US-09-375-257-3
16	97	68.3	23	1	US-08-333-565-10
17	97	68.3	23	2	US-08-661-479-10
18	97	68.3	59	2	US-08-733-505A-55
19	97	68.3	59	2	US-08-733-505A-56
20	97	68.3	59	2	US-08-733-505A-57
21	97	68.3	59	2	US-08-733-505A-58
22	81	57.0	16	1	US-08-333-565-26
23	81	57.0	16	2	US-08-661-479-26
24	56	39.4	11	2	US-08-733-505A-34
25	56	39.4	11	2	US-08-706-741B-69
26	56	39.4	11	2	US-08-924-695A-69
27	51	35.9	1125	4	US-09-252-991A-18729

28	48	33.8	626	4	US-09-252-991A-26276	Sequence 26276, A
29	48	33.8	1064	4	US-09-252-991A-17508	Sequence 17508, A
30	47	33.1	467	4	US-09-252-991A-18296	Sequence 18296, A
31	46	32.4	347	2	US-08-379-556A-2	Sequence 2, Appli
32	44	31.0	66	2	US-08-867-087B-40	Sequence 40, Appli
33	44	31.0	124	3	US-08-591-632-56	Sequence 56, Appli
34	44	31.0	124	4	US-09-611-451-56	Sequence 56, Appli
35	44	31.0	371	4	US-09-252-991A-20008	Sequence 20008, A
36	44	31.0	587	3	US-09-147-923-2	Sequence 2, Appli
37	44	31.0	724	4	US-09-328-352-7710	Sequence 7710, A
38	43.5	30.6	529	4	US-09-252-991A-22964	Sequence 22964, A
39	43.5	30.6	1047	4	US-09-198-452A-1058	Sequence 1058, Ap
40	43	30.3	125	4	US-09-328-352-7449	Sequence 7449, Ap
41	43	30.3	284	4	US-09-328-352-6559	Sequence 6559, Ap
42	43	30.3	2008	4	US-09-091-501B-8	Sequence 8, Appli
43	43	30.3	3433	4	US-09-091-501B-10	Sequence 10, Appli
44	42.5	29.9	460	4	US-09-252-991A-23461	Sequence 23461, A
45	42	29.6	28	1	US-08-261-660A-17	Sequence 17, Appli

ALIGNMENTS

RESULT 1
; Sequence 2, Application US/08665617
; Patent No. 5663316
; GENERAL INFORMATION:
; APPLICANT: Xudong, Yin
; TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Saliwanchik & Saliwanchik
; STREET: 2421 N.W. 41st Street, Suite A-1
; CITY: Gainesville
; STATE: Florida
; COUNTRY: USA
; ZIP: 32606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/665,617
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Saliwanchik, David R.
; REGISTRATION NUMBER: 31,794
; REFERENCE/DOCKET NUMBER: CL-8
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (352) 375-8100
; TELEFAX: (352) 372-5800
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 166 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-665-617-2

Query Match 96.5%; Score 137; DB 1; Length 166;
Best Local Similarity 96.3%; Pred. No. 53e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRMSDFVDFKGL 27
DB 101 NLWAAQYGRGLRMSDFVDFKGL 127

RESULT 2

```
US-08-717-123-2
; Sequence 2, Application US/08717123
; Patent No. 5965703
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
; TITLE OF INVENTION: Acids and Methods of Use
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/717.123
; FILING DATE: 20-SEP-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-ID 1929
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-717-123-2

Query Match 96.5%; Score 137; DB 2; Length 168;
Best Local Similarity 96.3%; Pred. No. 5-4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLNAAQYGRRLRMSEDFVDSFKKGL 27
Db 103 NLNAAQYGRRLRMSEDFVDSFKKGL 129

RESULT 3
US-08-985-335-1
; Sequence 1, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
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US-08-985-335-1
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 338673
US-08-985-335-1

Query Match 96.5%; Score 137; DB 3; Length 168;
Best Local Similarity 96.3%; Pred. No. 5-4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLNAAQYGRRLRMSEDFVDSFKKGL 27
Db 103 NLNAAQYGRRLRMSEDFVDSFKKGL 129

RESULT 4
US-08-985-335-7
; Sequence 7, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
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SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 1683637
US-09-985-335-7

Query Match 96.5%; Score 137; DB 3; Length 168;
Best Local Similarity 96.3%; Pred. No. 5.4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRELRMSDFVDSFKKGL 27
||||| |

Db 103 NLWAAQYGRELRMSDFVDSFKKGL 129

RESULT 5
US-09-410-372-1
Sequence 1, Application US/09410372
Patent No. 6281334
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/410,372
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/985,335
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0421 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: SYNORAB01
CLONE: 358673
US-09-410-372-1

Query Match 96.5%; Score 137; DB 3; Length 168;
Best Local Similarity 96.3%; Pred. No. 5.4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRELRMSDFVDSFKKGL 27

Db 103 NLWAAQYGRELRMSDFVDSFKKGL 129
||||| |

RESULT 6
US-09-410-372-7
Sequence 7, Application US/09410372
Patent No. 6281334
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/410,372
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/985,335
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0421 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 1683637
US-09-410-372-7

Query Match 96.5%; Score 137; DB 3; Length 168;
Best Local Similarity 96.3%; Pred. No. 5.4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRELRMSDFVDSFKKGL 27
||||| |

Db 103 NLWAAQYGRELRMSDFVDSFKKGL 129

RESULT 7
US-09-375-257-2
Sequence 2, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
FILE OF INVENTION: ACIDS AND METHODS OF USE
FILE REFERENCE: 480140, 428D1
CURRENT APPLICATION NUMBER: US/09/375,257


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; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-375-257-2

Query Match          96.5%; Score 137; DB 4; Length 168;
Best Local Similarity 96.3%; Pred. No. 5.4e-14;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFVDSFKKL 27
Db 103 NLWAAQRYGRELRRMSDEFVDSFKKL 129

RESULT 8
US-08-333-565-2
; Sequence 2, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "Deduced amino acid sequence
; OF mouse BAD."
US-08-333-565-2

Query Match          76.8%; Score 109; DB 1; Length 204;
Best Local Similarity 87.5%; Pred. No. 1.7e-09;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFVDSFK 24
Db 140 NLWAAQRYGRELRRMSDEFVDSFK 163

RESULT 10
US-08-733-505A-1
; Sequence 1, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
```

ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-733-505A-1

Query Match 76.8%; Score 109; DB 2; Length 204;
Best Local Similarity 87.5%; Pred. No. 1.7e-09;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRRLRMSDFVDSFK 24
||||| ||||||| |||
DB 140 NLWAAQYGRRLRMSDFEGSFK 163

RESULT 11
US-08-733-505A-12
Sequence 12, Application US/08733505A
Patent No. 5856445
GENERAL INFORMATION:
APPLICANT: KORSMEYER, STANLEY J.
TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
NUMBER OF SEQUENCES: 60
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOWELL & HAFERKAMP, L.C.
CITY: ST. LOUIS
STATE: MISSOURI
COUNTRY: USA
ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear

MOLECULE TYPE: peptide
US-08-733-505A-12
Query Match 76.8%; Score 109; DB 2; Length 204;
Best Local Similarity 87.5%; Pred. No. 1.7e-09;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 NLWAAQYGRRLRMSDFVDSFK 24
||||| ||||||| |||
DB 140 NLWAAQYGRRLRMSDFEGSFK 163
RESULT 12
US-08-733-505A-13
Sequence 13, Application US/08733505A
Patent No. 5856445
GENERAL INFORMATION:
APPLICANT: KORSMEYER, STANLEY J.
TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
NUMBER OF SEQUENCES: 60
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOWELL & HAFERKAMP, L.C.
CITY: ST. LOUIS
STATE: MISSOURI
COUNTRY: USA
ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-13

Query Match 76.8%; Score 109; DB 2; Length 204;
Best Local Similarity 87.5%; Pred. No. 1.7e-09;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRRLRMSDFVDSFK 24
||||| ||||||| |||
DB 140 NLWAAQYGRRLRMSDFEGSFK 163

RESULT 13
US-08-733-505A-14
Sequence 14, Application US/08733505A
Patent No. 5856445
GENERAL INFORMATION:
APPLICANT: KORSMEYER, STANLEY J.
TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
NUMBER OF SEQUENCES: 60
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOWELL & HAFERKAMP, L.C.

STREET: 7733 FORSYTH BLVD., SUITE 1400
CITY: ST. LOUIS
STATE: MISSOURI
COUNTRY: USA
ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-14
Query Match 76.8%; Score 109; DB 2; Length 204;
Best Local Similarity 87.5%; Pred. No. 1.7e-09;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 NLWAAQYRGRLRRMSDFVDSFK 24
Db 140 NLWAAQYRGRLRRMSDFEGSFK 163
RESULT 14
US-08-717-123-3
Sequence 3, Application US/08717123
Patent No. 5965703
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
TITLE OF INVENTION: Acids and Methods of Use
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell and Flores
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: United States
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/717,123
FILING DATE: 20-SEP-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-ID 1929
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 3:

SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
TOPOLOGY: linear
US-08-717-123-3
Query Match 74.6%; Score 106; DB 2; Length 204;
Best Local Similarity 83.3%; Pred. No. 5e-09;
Matches 20; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
QY 1 NLWAAQYRGRLRRMSDFVDSFK 24
Db 140 NLWAAQYRGRLRRMTDEFGSFK 163
RESULT 15
US-09-375-257-3
Sequence 3, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
TITLE OF INVENTION: ACIDS AND METHODS OF USE
FILE REFERENCE: 480140.428D1
CURRENT APPLICATION NUMBER: US/09/375,257
CURRENT FILING DATE: 1999-08-16
NUMBER OF SEQ ID NOS: 15
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 3
LENGTH: 204
TYPE: PRT
ORGANISM: Mus musculus
US-09-375-257-3
Query Match 74.6%; Score 106; DB 4; Length 204;
Best Local Similarity 83.3%; Pred. No. 5e-09;
Matches 20; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
QY 1 NLWAAQYRGRLRRMSDFVDSFK 24
Db 140 NLWAAQYRGRLRRMTDEFGSFK 163
Search completed: September 23, 2003, 09:45:40
Job time : 32 secs

Result No.	Score	Query		DB	ID	Description
		Match	Length			
1	137	96.5	168	9	US-09-922-378-2	Sequence 2, Appli
2	137	96.5	168	9	US-09-894-657-1	Sequence 1, Appli
3	137	96.5	168	9	US-09-894-657-7	Sequence 2, Appli
4	137	96.5	168	12	US-10-209-967-2	Sequence 7, Appli
5	137	96.5	168	14	US-10-066-179-2	Sequence 2, Appli
6	127	89.4	215	15	US-10-059-261-258	Sequence 258, App
7	109	76.8	204	12	US-10-209-967-4	Sequence 4, Appli
8	106	74.6	204	9	US-09-922-378-3	Sequence 3, Appli
9	106	74.6	204	14	US-10-066-179-3	Sequence 3, Appli
10	57	40.1	373	15	US-10-374-108A-147	Sequence 147, App
11	48	33.8	373	12	US-10-168-780-2	Sequence 2, Appli
12	47	33.1	350	11	US-09-940-244-394	Sequence 394, App
13	46.5	32.7	1053	9	US-09-841-132-583	Sequence 583, App
14	46.5	32.7	1053	14	US-10-007-693-97	Sequence 97, Appl
15	46	32.4	1053	15	US-10-092-750-1	Sequence 1, Appli

RESULT 2
US-09-894-657-1
Sequence 1, Application US/09894657
; Patent No. US2002009859A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; yue, Henry
; Lal, Preeti
; Shah, Purvi
;

REF. 94504
COMPUTER READABLE FORM:
MEDIUM TYPE: Disk
COMPUTER: IBM Comp

RESULT 5
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.

; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-066-179-2

Query Match 96.5%; Score 137; DB 14; Length 168;
Best Local Similarity 96.3%; Pred. No. 1.2e-12;
Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRRMSDEFVDSFKGL 27
DB 103 NLWAAQYGRGLRRMSDEFVDSFKGL 129

RESULT 6

US-10-059-261-258
; Sequence 258, Application US/10059261
; Publication No. US20030077826A1
; GENERAL INFORMATION:
; APPLICANT: EDELMAN, LENA
; APPLICANT: JACOTOT, JEAN-PAUL
; APPLICANT: BRIAND, DANIEL FRANCOIS
; TITLE OF INVENTION: CHIMERIC MOLECULES CONTAINING A MODULE ABLE TO TARGET
; TITLE OF INVENTION: SPECIFIC CELLS AND A MODULE REGULATING THE APOPTOTIC
; TITLE OF INVENTION: FUNCTION OF THE PERMEABILITY TRANSITION PORE COMPLEX
; TITLE OF INVENTION: (PTPC)
; FILE REFERENCE: 03495.0216
; CURRENT APPLICATION NUMBER: US/10/059,261
; PRIOR FILING DATE: 2002-08-29
; PRIOR APPLICATION NUMBER: 60/255,594
; PRIOR FILING DATE: 2001-02-02
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 258
; LENGTH: 25
; TYPE: PRT
; ORGANISM: Unknown Organism
; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: TOX peptide
US-10-059-261-258

Query Match 89.4%; Score 127; DB 15; Length 25;
Best Local Similarity 96.0%; Pred. No. 4.6e-12;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRRMSDEFVDSFKK 25
DB 1 NLWAAQYGRGLRRMSDEFVDSFKK 25

RESULT 7

US-10-209-967-4
; Sequence 4, Application US/10209967
; Publication No. US20030171279A1
; GENERAL INFORMATION:
; APPLICANT: MUNGER, JOSHUA
; APPLICANT: ROIZMAN, BERNARD
; TITLE OF INVENTION: METHODS AND COMPOSITION CONCERNING HERPESVIRUS US3 AND
; TITLE OF INVENTION: RAD-INVOLVED APOPTOSIS
; FILE REFERENCE: ARCD:380US
; CURRENT APPLICATION NUMBER: US/10/209,967
; CURRENT FILING DATE: 2002-07-31
; PRIOR APPLICATION NUMBER: 60/308,929
; PRIOR FILING DATE: 2001-07-31

; NUMBER OF SEQ ID NOS: 25
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 4
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-209-967-4

Query Match 76.8%; Score 109; DB 12; Length 204;
Best Local Similarity 87.5%; Pred. No. 2.1e-08;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRRMSDEFVDSFK 24
DB 140 NLWAAQYGRGLRRMSDEFVDSFK 163

RESULT 8

US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 74.6%; Score 106; DB 9; Length 204;
Best Local Similarity 83.3%; Pred. No. 5.8e-08;
Matches 20; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRRMSDEFVDSFK 24
DB 140 NLWAAQYGRGLRRMTDEFVDSFK 163

RESULT 9

US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-066-179-3

Query Match 74.6%; Score 106; DB 14; Length 204;
Best Local Similarity 83.3%; Pred. No. 5.8e-08;
Matches 20; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRGLRRMSDEFVDSFK 24
DB 140 NLWAAQYGRGLRRMTDEFVDSFK 163

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RESULT 10
US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US20030068652A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECIFIC,
; TITLE OF INVENTION: CONTEXT-INDEPENDENT ANTIBODIES COUPLED WITH DATABASE SEARCHING
; FILE REFERENCE: CST-138 CIP3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147
Query Match 40.1%; Score 57; DB 15; Length 15;
Best Local Similarity 85.7%; Pred. No. 0.063; 2; Indels 0; Gaps 0;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 9 GRELRMSDEFVDS 22
Db 1 GRELRMSDEFGS 14

RESULT 11
US-10-168-780-2
; Sequence 2, Application US/10168780
; Publication No. US20030172405A1
; GENERAL INFORMATION:
; APPLICANT: TANAKA, Hitoshi
; APPLICANT: KAYANO, Toshiaki
; APPLICANT: YANO, Masahiro
; APPLICANT: MATSUOKA, Makoto
; APPLICANT: KOBAYASHI, Masatomo
; TITLE OF INVENTION: GIBBERELLIN 3BETA-HYDROXYLASE GENES OF RICE AND USES THEREOF
; FILE REFERENCE: SH2-004US
; CURRENT APPLICATION NUMBER: US/10/168,780
; CURRENT FILING DATE: 2002-06-20
; PRIOR APPLICATION NUMBER: PCT/JP00/09037
; PRIOR FILING DATE: 2000-12-20
; PRIOR APPLICATION NUMBER: JP 11/361608
; PRIOR FILING DATE: 1999-12-20
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: PatentIn ver. 2.0
; SEQ ID NO 2
; LENGTH: 373
; TYPE: PRT
; ORGANISM: Oryza sativa
US-10-168-780-2
Query Match 33.8%; Score 48; DB 12; Length 373;
Best Local Similarity 36.4%; Pred. No. 45;
Matches 8; Conservative 8; Mismatches 6; Indels 6; Gaps 0;

QY 6 QEYGRRLRMSDEFVDSFKGL 27
Db 162 EEFKEMRLADELLRLFLRAL 183

RESULT 12
US-09-940-244-394
; Sequence 394, Application US/09940244
; Publication No. US20030044796A1
; GENERAL INFORMATION:
; APPLICANT: Neri, Bruce P.
; APPLICANT: Hall, Jeff G.
; APPLICANT: Lyamichev, Victor
; APPLICANT: Smith, Lloyd M.
; TITLE OF INVENTION: Reactions on Dendrimers
; FILE REFERENCE: FORS-06478
; CURRENT APPLICATION NUMBER: US/09/940,244
; CURRENT FILING DATE: 2002-05-06
; NUMBER OF SEQ ID NOS: 422
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 394
; LENGTH: 350
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-09-940-244-394
Query Match 33.1%; Score 47; DB 11; Length 350;
Best Local Similarity 40.9%; Pred. No. 58;
Matches 9; Conservative 5; Mismatches 8; Indels 0; Gaps 0;

QY 4 AAQYGRRLRMSDEFVDSFKK 25
Db 119 BARYAQAAARLIDENVEDAKK 140

RESULT 13
US-09-841-132-583
; Sequence 583, Application US/09841132
; Patent No. US20020061848A1
; GENERAL INFORMATION:
; APPLICANT: Bhatia, Ajay
; APPLICANT: Skeiky, Yasir A.W.
; APPLICANT: Probst, Peter
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATMENT AND
; TITLE OF INVENTION: DIAGNOSIS OF CHLAMYDIAL INFECTION
; FILE REFERENCE: 210121.469C8
; CURRENT APPLICATION NUMBER: US/09/841,132
; CURRENT FILING DATE: 2001-04-23
; NUMBER OF SEQ ID NOS: 599
; SOFTWARE: FastSeq for Windows Version 3.0/4.0
; SEQ ID NO 583
; LENGTH: 1053
; TYPE: PRT
; ORGANISM: C. Trachomatis D serovar
US-09-841-132-583
Query Match 32.7%; Score 46.5; DB 9; Length 1053;
Best Local Similarity 40.7%; Pred. No. 2.3e+02;
Matches 11; Conservative 7; Mismatches 6; Indels 3; Gaps 1;

QY 1 NLWAAQYGRRLRMSDEFVDSFKGL 27
Db 969 NLVLAQPDGKKLSNM--YLTAWKKGL 992

RESULT 14
US-10-007-693-97
; Sequence 97, Application US/10007693
; Publication No. US20020146776A1
; GENERAL INFORMATION:
; APPLICANT: Bhatia, Ajay
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QY 6 QEYGRRLRMSDEFVDSFKGL 27
Db 162 EEFKEMRLADELLRLFLRAL 183

RESULT 12
US-09-940-244-394
; Sequence 394, Application US/09940244
; Publication No. US20030044796A1
; GENERAL INFORMATION:
; APPLICANT: Neri, Bruce P.
; APPLICANT: Hall, Jeff G.
; APPLICANT: Lyamichev, Victor
; APPLICANT: Smith, Lloyd M.
; TITLE OF INVENTION: Reactions on Dendrimers
; FILE REFERENCE: FORS-06478
; CURRENT APPLICATION NUMBER: US/09/940,244
; CURRENT FILING DATE: 2002-05-06
; NUMBER OF SEQ ID NOS: 422
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 394
; LENGTH: 350
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-09-940-244-394
Query Match 33.1%; Score 47; DB 11; Length 350;
Best Local Similarity 40.9%; Pred. No. 58;
Matches 9; Conservative 5; Mismatches 8; Indels 0; Gaps 0;

QY 4 AAQYGRRLRMSDEFVDSFKK 25
Db 119 BARYAQAAARLIDENVEDAKK 140

RESULT 13
US-09-841-132-583
; Sequence 583, Application US/09841132
; Patent No. US20020061848A1
; GENERAL INFORMATION:
; APPLICANT: Bhatia, Ajay
; APPLICANT: Skeiky, Yasir A.W.
; APPLICANT: Probst, Peter
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATMENT AND
; TITLE OF INVENTION: DIAGNOSIS OF CHLAMYDIAL INFECTION
; FILE REFERENCE: 210121.469C8
; CURRENT APPLICATION NUMBER: US/09/841,132
; CURRENT FILING DATE: 2001-04-23
; NUMBER OF SEQ ID NOS: 599
; SOFTWARE: FastSeq for Windows Version 3.0/4.0
; SEQ ID NO 583
; LENGTH: 1053
; TYPE: PRT
; ORGANISM: C. Trachomatis D serovar
US-09-841-132-583
Query Match 32.7%; Score 46.5; DB 9; Length 1053;
Best Local Similarity 40.7%; Pred. No. 2.3e+02;
Matches 11; Conservative 7; Mismatches 6; Indels 3; Gaps 1;

QY 1 NLWAAQYGRRLRMSDEFVDSFKGL 27
Db 969 NLVLAQPDGKKLSNM--YLTAWKKGL 992

RESULT 14
US-10-007-693-97
; Sequence 97, Application US/10007693
; Publication No. US20020146776A1
; GENERAL INFORMATION:
; APPLICANT: Bhatia, Ajay
```

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; APPLICANT: Probst, Peter
; TITLE OF INVENTION: COMPOUNDS AND METHODS FOR TREATMENT
; FILE OF INVENTION: AND DIAGNOSIS OF CHLAMYDIAL INFECTION
; FILE REFERENCE: 210121.515C2
; CURRENT APPLICATION NUMBER: US/10/007,693
; CURRENT FILING DATE: 2001-12-05
; NUMBER OF SEQ ID NOS: 157
; SEQ ID NO 97
; LENGTH: 1053
; TYPE: PRT
; ORGANISM: Chlamydia trachomatis serovar D
US-10-007-693-97
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Query Match 32.7%; Score 46.5; DB 14; Length 1053;
Best Local Similarity 40.7%; Pred. No. 2.3e+02;
Matches 11; Conservative 7; Mismatches 6; Indels 3; Gaps 1;
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```
QY 1 NLWAAQEGYGRELRMSDEFVDFKGL 27
   ||:| |::| |:::||||
Db 969 NYLAQPDGKKLSNM---YLTANKGL 992
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RESULT 15

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US-10-092-750-1
; Sequence 1, Application US/10092750
; Publication No. US20030032157A1
; GENERAL INFORMATION:
; APPLICANT: Hammond, Philip W.
; APPLICANT: Alpin, Julia
; APPLICANT: Wright, Martin C.
; TITLE OF INVENTION: Polypeptides Interactive with BCL-XL
; FILE REFERENCE: 50036/050002
; CURRENT APPLICATION NUMBER: US/10/092,750
; CURRENT FILING DATE: 2002-03-07
; PRIOR APPLICATION NUMBER: US 60/274,526
; PRIOR FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 253
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 35
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-092-750-1
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Query Match 32.4%; Score 46; DB 15; Length 35;
Best Local Similarity 55.6%; Pred. No. 6.8;
Matches 10; Conservative 2; Mismatches 2; Indels 4; Gaps 1;
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QY 2 LWAAQEGYGRELRMSDEF 19
   :| | |:::||||
Db 15 IWTAQ---ELRRIGDEF 28
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Search completed: September 23, 2003, 10:03:35
Job time : 390 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 23, 2003, 09:43:16 ; Search time 22 Seconds
(without alignments)
57.715 Million cell updates/sec

Title: US-09-544-664B-2

Perfect score: 142

Sequence: 1 NLWAAQEGYRELRLMSDFVDSFKKGL 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Query %	ID	Description
1	137	96.5		1 BAD_HUMAN	Q92934 homo sapien
2	109	76.8		1 BAD_MOUSE	Q61337 mus musculus
3	109	76.8		1 BAD_RAT	Q35147 rattus norv
4	51	35.9		1 HT2A_HUMAN	Q13049 homo sapien
5	50.5	35.6		1 MTBR_BPH1	P09915 bacterioph
6	50.5	35.6		1 PROG_THTN	Q8r943 thermocae
7	50	35.2		1 AROG_YEAST	P32449 saccharomyc
8	49.5	34.9		1 BXDI_DROME	Q9veb3 drosophila
9	49	34.5		1 HXK_DEBOC	P50506 debaryomyce
10	48	33.8		1 AROG_CANAL	P79023 candida alb
11	48	33.8		1 VP5_BT2A	P30209 bluetongue
12	48	33.8		1 SYK_THEMA	O33925 thermotoga
13	47.5	33.5		1 PCK_RHISN	P43086 rhizobium s
14	47.5	33.5		1 VGI3_BPM2	O64206 mycobacteri
15	47	33.1		1 RSPD_BORBU	P52323 borrelia bu
16	47	33.1		1 CS05_MOUSE	Q8k2h3 mus musculu
17	46.5	32.7		1 IPYR_MYCPE	O8evw3 mycoplasma
18	46.5	32.7		1 T591_AQUAE	O6f853 aquifex aeo
19	46.5	32.7		1 RRL1_CHLMO	Q9p193 chlamydia m
20	46.5	32.7		1 RRL1_CHLTR	O43521 homo sapien
21	46	32.4		1 B1M_HUMAN	O43521 homo sapien
22	46	32.4		1 MRAW_RICPR	Q92cy2 rickettsia
23	46	32.4		1 FLS_PETHY	O07512 petunia hyb
24	46	32.4		1 TNA4_AERPE	Q9yc12 aeropyrum p
25	46	32.4		1 F5P_BTIVA	P12436 bluetongue
26	46	32.4		1 STUL_YEAST	P38198 saccharomyc
27	45.5	32.0		1 XYAD_BACSU	P42103 bacillus su
28	45.5	32.0		1 PPCK_BACHD	Q9K7g7 bacillus su
29	45.5	32.0		1 XYBT_BACSU	P37484 bacillus su
30	45	31.7		1 SVET_METUA	Q56443 methanococc
31	45	31.7		1 SCVA_HABIN	P43803 haemophilus
32	44.5	31.3		1 ISC1_ARCFU	O30052 archaeoglob
33	44.5	31.3		1 DPOA_LEIDO	O00874 leishmania

34 44 31.0 125 1 VSIS_REOVJ P32868 reovirus (t
35 44 31.0 232 1 A23_ARATH P35632 arabidopsis
36 44 31.0 261 1 YGF_ECOLI P2045 escherichia
37 44 31.0 310 1 ACP2_ENTHI P36185 entamoeba h
38 44 31.0 311 1 ALSB_ECOLI P39265 escherichia
39 44 31.0 315 1 CPP2_ENTHI Q01958 entamoeba h
40 44 31.0 351 1 E2BB_RABIT Q28690 oryctolagus
41 44 31.0 422 1 PANL_HUMAN Q96rd7 homo sapien
42 44 31.0 453 1 RMOC_PSAE Q914u3 pseudomonas
43 44 31.0 534 1 CN9A_MOUSE O70628 mus musculu
44 44 31.0 661 1 HT31_ARATH Q04996 arabidopsis
45 44 31.0 704 1 KPCL_CABEL P34722 caenorhabdi

ALIGNMENTS

RESULT 1
BAD_HUMAN STANDARD; PRT; 168 AA.
AC Q92934; Q14803;
DT 01-NOV-1997 (Rel. 35, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-
XL/Bcl-2 associated death promoter) (BCL2-like 8 protein).
GN BAD OR BCL2 OR BCL2L6.
OS Homo sapiens (Human).
OC Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Yin D.X., Li Z., Huang B., Chen S., Zhou H.;
RT "A human protein that interacts with Bcl-2 and have homology to mouse
RT BAD.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.
RX MEDLINE=97083574; PubMed=8929532;
RA Wang H.-G., Rapp U.R., Reed J.C.;
RT "Bcl-2 targets the protein kinase Raf-1 to mitochondria.";
RN [3]
RP SEQUENCE FROM N.A.
RA Takayama S., Reed J.C.;
RT Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A., AND DIMERIZATION.
RX MEDLINE=98049554; PubMed=9388232;
RA Ottile S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,
RT Chang S., Weeks S., Fritz L.C., Oltersdorf T.;
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyar S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,

Ra Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 Ra Butterfield J.S.N., Krzywinski M.I., Skalska U., Smalilus D.E.,
 Ra Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
 Rt "Generation and initial analysis of more than 15,000 full-length
 Rt human and mouse cDNA sequences";
 Rl Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 Rn [6]
 Rn STRUCTURE BY NMR OF 103-127.
 Rx MEDLINE=21073561; PubMed=11206074;
 Ra Petros A.M., Nettesheim D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
 Ra Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
 Ra Fesik S.W.;
 Rt "Rationale for Bcl-xL/Bad peptide complex formation from structure,
 Rt mutagenesis, and biophysical studies";
 Rl Protein Sci. 9:2528-2534(2000).
 Cc -!- FUNCTION: Promotes cell death. Successfully competes for the
 Cc binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 Cc of heterodimerization of these proteins with BAX. Can reverse the
 Cc death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 Cc similarity). Appears to act as a link between growth factor
 Cc receptor signaling and the apoptotic pathways.
 Cc -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 Cc x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 Cc The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
 Cc similarity).
 Cc -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 Cc phosphorylation, locates to the cytoplasm.
 Cc -!- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
 Cc -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 Cc BAX for their pro-apoptotic activity and for their interaction
 Cc with anti-apoptotic members of the Bcl-2 family.
 Cc -!- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
 Cc Subsequent phosphorylation on Ser-99 promotes heterodimerization
 Cc with 14-3-3 proteins. This interaction then facilitates the
 Cc phosphorylation at Ser-118, a site within the BH3 domain, leading
 Cc to the release of Bcl-x(L) and the promotion of cell survival.
 Cc Ser-99 is the major site of AKT/PKB phosphorylation, Ser-118 the
 Cc major site of protein kinase A (CAPK) phosphorylation (By
 Cc similarity).
 Cc -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
 Cc -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 Cc -!- CAUTION: Ref1 sequence differs from that shown due to frameshifts
 Cc in position 64 and 91.
 Cc -----
 Cc This SWISS-PROT entry is copyright. It is produced through a collaboration
 Cc between the Swiss Institute of Bioinformatics and the EMBL outstation -
 Cc the European Bioinformatics Institute. There are no restrictions on its
 Cc use by non-profit institutions as long as its content is in no way
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 Cc or send an email to license@isb-sib.ch).
 Cc -----
 Dr EMBL; U66879; AAB36516.1; ALT_FRAME.
 Dr EMBL; AF021792; AAB72092.1; -
 Dr EMBL; AF031523; AAB86124.1; -
 Dr EMBL; BC001901; AAB01901.1; -
 Dr PDB; 1G5J; 07-FEB-01.
 Dr Genew; HGNC:936; BAD.
 Dr MTM; 603167; -
 Dr GO; GO:0005737; C:cytoplasm; NAS.
 Dr GO; GO:0005741; C:mitochondrial outer membrane; NAS.
 Dr GO; GO:0005515; F:protein binding activity; NAS.
 Dr GO; GO:0008632; P:apoptotic program; TAS.
 Dr GO; GO:0006917; P:induction of apoptosis; NAS.
 Dr InterPro; IPR000712; Bcl2.BH
 Dr PROSITE; PS01259; BH3; FALSE_NEG.
 Kw Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
 Ft DOMAIN 110 124 BH3.
 Ft MOD_RES 75 75 PHOSPHORYLATION (BY PKA AND PKB) (BY
 Ft SIMILARITY).
 Ft MOD_RES 99 99 PHOSPHORYLATION (BY PKA AND PKB) (BY
 Ft SIMILARITY).
 Ft MOD_RES 118 118 PHOSPHORYLATION (BY PKA AND PKB) (BY
 Ft SIMILARITY).

FT VARIANT 107 107 SIMILARITY).
 FT A -> S (in dbSNP:3729933).
 FT /FTID=VAR_015380.
 FT HELIX 106 121
 SQ SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEB3241 CRC64;
 Query Match 96.5%; Score 137; DB-1; Length 168;
 Best Local Similarity 96.3%; Pred. No. 8.2e-13;
 Matches 26; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 NLWAAQYGHRLRMSDFVDSFKKGL 27
 Db 103 NLWAAQYGHRLRMSDFVDSFKKGL 129
 RESULT 2
 BAD_MOUSE STANDARD; PRT; 204 AA.
 ID BAD_MOUSE Q61337;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 26-FEB-2003 (Rel. 41, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component
 DE 6) (Bcl-xL/Bcl-2 associated death promoter).
 GN BAD OR BCC6.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain, and Thymus;
 RX MEDLINE=95136361; PubMed=7834748;
 RA Yang E., Zha J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;
 RT "Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and
 RT promotes cell death.";
 RL Cell 80:285-291(1995).
 RN [2]
 RP PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.
 RX MEDLINE=98022383; PubMed=9381178;
 RA Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;
 RT "Interleukin-3-induced phosphorylation of BAD through the protein
 RT kinase Akt.";
 RL Science 278:687-689(1997).
 RN [3]
 RP MUTAGENESIS OF SERINE RESIDUES.
 RX MEDLINE=20403302; PubMed=10949026;
 RA Datta S.R., Katsov A., Hu L., Petros A., Fesik S.W., Yaffe M.B.,
 RA Greenberg M.E.;
 RT "14-3-3 proteins and survival kinases cooperate to inactivate BAD by
 RT BH3 domain phosphorylation.";
 RL Mol. Cell 6:41-51(2000).
 Cc -!- FUNCTION: Promotes cell death. Successfully competes for the
 Cc binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 Cc of heterodimerization of these proteins with BAX. Can reverse the
 Cc death repressor activity of Bcl-x(L), but not that of Bcl-2.
 Cc Appears to act as a link between growth factor receptor signaling
 Cc and the apoptotic pathways.
 Cc -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 Cc x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 Cc The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.
 Cc -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 Cc phosphorylation, locates to the cytoplasm.
 Cc -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 Cc BAX for their pro-apoptotic activity and for their interaction
 Cc with anti-apoptotic members of the Bcl-2 family.
 Cc -!- PTM: Phosphorylated on Ser-112 in response to survival stimuli.
 Cc Subsequent phosphorylation on Ser-136 promotes heterodimerization
 Cc with 14-3-3 proteins. This interaction then facilitates the
 Cc phosphorylation at Ser-155, a site within the BH3 domain, leading
 Cc to the release of Bcl-x(L) and the promotion of cell survival.
 Cc Ser-136 is the major site of AKT/PKB phosphorylation, Ser-155 the
 Cc major site of protein kinase A (CAPK) phosphorylation.


```
Best Local Similarity 87.5%; Pred. No. 1.2e-08;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 NLWAAQYGRRLRMSDFVDSFK 24
    ||||| ||||| ||||| |||||
DB 141 NLWAAQYGRRLRMSDFVDSFK 164

RESULT 4
ID HT2A_HUMAN STANDARD; PRT; 653 AA.
AC Q13049; Q9NQPB;
DT 01-NOV-1997 (Rel. 35, Created)
DT 15-FEB-2003 (Rel. 41, Last sequence update)
DT 15-FEB-2003 (Rel. 42, Last annotation update)
DE Zinc-finger protein HT2A (72 kDa Tat-interacting protein) (Tripartite
DE motif-containing protein 32).
GN TRIM32 OR HT2A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=95297135; PubMed=7778269;
RA Fridell R.A., Harding L.S., Bogerd H.P., Cullen B.R.;
RT "Identification of a novel human zinc finger protein that
RT specifically interacts with the activation domain of lentiviral Tat
RT proteins."
RL Virology 209:347-357(1995).
RN [2]
RP SEQUENCE FROM N.A.
RA Sehra H.;
RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Skib;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausberg R.D., Collins F.S., Wagner K.H., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan K., Moore T., Max S.I., Wang J., Hsieh F.,
RA Dlatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raba S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Kryzyski M.I., Skalska U., Smalhus D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- FUNCTION: MAY PLAY A SIGNIFICANT ROLE IN MEDIATING THE BIOLOGICAL
CC ACTIVITY OF THE HIV-1 TAT PROTEIN IN VIVO. BINDS SPECIFICALLY TO
CC THE ACTIVATION DOMAIN OF HIV-1 TAT AND CAN ALSO INTERACT WITH THE
CC HIV-2 AND ELAV TAT PROTEINS IN VIVO.
CC -!- SUBCELLULAR LOCATION: Nuclear.
CC -!- TISSUE SPECIFICITY: SPLEEN, THYMUS, PROSTRATE, TESTIS, OVARY,
CC INTESTINE AND COLON.
CC -!- SIMILARITY: Contains 1 RING-type zinc finger.
CC -!- SIMILARITY: Contains 1 B box-type zinc finger.
CC -----
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CC -----
DR EMBL; U18543; AAA6474.1; -.
DR EMBL; AL133284; CAB92723.1; -.
DR EMBL; BC003154; AAH03154.1; -.
DR HSSP; P29590; 1BOR.
DR Genew; HGNC:16380; TRIM32.
DR MIM; 602230; -.
DR GO; GO:0005634; C:nucleus; TAS.
DR GO; GO:0003713; F:transcription co-activator activity; TAS.
DR InterPro; IPR001258; NHL.
DR InterPro; IPR000315; Znf_Box.
DR InterPro; IPR001841; Znf_ring.
DR Pfam; PF01436; NHL; 5.
DR Pfam; PF00643; zf-B_Box; 1.
DR Pfam; PF00097; zf-C3HC4; 1.
DR SMART; SM00336; BBOX; 1.
DR SMART; SM00184; RING; 1.
DR PROSITE; PS00119; ZF_BOX; 1.
DR PROSITE; PS00518; ZF_RING_1; 1.
DR PROSITE; PS00089; ZF_RING_2; 1.
KW Zinc-finger; Nuclear protein.
FT DOMAIN 2 6 POLY-ALA.
FT ZN_FING 20 65 RING-TYPE.
FT ZN_FING 103 133 B_BOX-TYPE.
FT CONFLICT 27 27 F -> I (IN REF. 1).
SQ SEQUENCE 653 AA; 71988 MW; D83B1595CA8378FD CRC64;

Query Match 35.9%; Score 51; DB 1; Length 653;
Best Local Similarity 57.9%; Pred. No. 11;
Matches 11; Conservative 1; Mismatches 7; Indels 0; Gaps 0;

QY 6 QEYGRRLRMSDFVDSFK 24
    |||| | ||: || | |
DB 186 QEYGRRLRMSDFVDSFK 204

RESULT 5
ID MTER_BPRH1 STANDARD; PRT; 503 AA.
AC P09915;
DT 01-MAR-1989 (Rel. 10; Created)
DT 01-NOV-1995 (Rel. 32; Last sequence update)
DT 16-OCT-2001 (Rel. 40; Last annotation update)
DE Modification methylase RhoIIsi (EC 2.1.1.73) (Cytosine-specific
DE methyltransferase RhoIIsi) (Ssu Flis) (M.RhoIIsi).
OS Bacteriophage rho-IIs.
OC Viruses; dsDNA viruses, no RNA stage; Caudovirales; Siphoviridae;
OC Lambda-like viruses.
OX NCBI_TaxID=10735;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87246516; PubMed=3109889;
RA Behrens B., Noyer-Weidner M., Pawlek B., Lauster R., Balganesch T.S.,
RA Trautner T.A.;
RT "Organization of multispecific DNA methyltransferases encoded by
RT temperate Bacillus subtilis phages."
RL EMBO J. 6:1137-1142(1987).
RN [2]
RP REVISION TO 476.
RA Trautner T.A.;
RL Submitted (SEP-1987) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: THIS ENZYME METHYLATES CYTOSINE WITHIN THE SEQUENCES
CC GACC AND GAGTCC.
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + DNA cytosine = S-
CC adenosyl-L-homocysteine + DNA 5-methylcytosine.
CC -!- SIMILARITY: BELONGS TO THE C5-METHYLTRANSFERASE FAMILY.
CC -----
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EMBL; X05242; CAA28869.1; -
 DR F1R; A28137; CTEPRH.
 DR HSP; P05102; GMYH.
 DR RERASE; 2835; M.RhollisI.
 DR InterPro; IPR001525; C5_DNA_meth.
 DR Pfam; PF00145; DNA_methylase; 1.
 DR PRINTS; PR00105; C5METTRFRASE.
 DR TIGRFAMS; TIGR00675; dcm; 1.
 DR PROSITE; PS00094; C5_MTASE_1; 1.
 DR PROSITE; PS00095; C5_MTASE_2; 1.
 KW Transferase; Methyltransferase; Restriction system.
 FT ACT SITE 78 78 BY SIMILARITY.
 SQ SEQUENCE 503 AA; 57129 MW; AAFB6FE01B8129E CRC64;

Query Match 35.6%; Score 50.5; DB 1; Length 503;
 Best Local Similarity 50.0%; Pred. No. 9.5;
 Matches 10; Conservative 5; Mismatches 4; Indels 1; Gaps 1;

QY 3 WAAQ-YGRELRMSDEVD 21
 |:|:|: |:|:|: |:|:|:|
 Db 207 WSAQDIVGRRLREILEYVD 226

RESULT 6

PPCK_THEN
 ID PPCK_THEN STANDARD; PRT; 521 AA.
 AC Q8R943;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Phosphoenolpyruvate carboxylase [ATP] (EC 4.1.1.49) (PEP
 DE carboxylase) (Phosphoenolpyruvate carboxylase) (PEPCK).
 GN PCKA OR TPE1783.
 OS Thermoanaerobacter tengcongensis.
 OC Bacteria; Firmicutes; Clostridia; Thermoanaerobacteriales;
 OC Thermoanaerobacteriaceae; Thermoanaerobacter.
 OX NCBI_TaxID=119072;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MB4 / JCM 11007;
 RX MEDLINE=21992816; PubMed=11997336;
 RA Bao Q., Tian Y., Li W., Xu Z., Xuan Z., Hu S., Dong W., Yang J.,
 RA Chen Y., Xue Y., Xu Y., Lai X., Huang L., Dong X., Ma Y., Ling L.,
 RA Tan H., Chen R., Wang J., Yu J., Yang H.;
 RT "A complete sequence of T. tengcongensis genome.";
 RL Genome Res. 12:689-700(2002).
 CC -!- CATALYTIC ACTIVITY: ATP + oxaloacetate = ADP + phosphoenolpyruvate
 CC + Co(2).
 CC -!- PATHWAY: Rate-limiting gluconeogenic enzyme.
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
 CC -!- SIMILARITY: Belongs to the phosphoenolpyruvate carboxylase [ATP]
 CC family.
 CC
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EMBL; AE013131; AA024977.1; -
 DR HAMAP; MF_00453; -; 1.
 DR InterPro; IPR001272; PEPCK_ATP.
 DR Pfam; PF01293; PEPCK_ATP; 1.
 DR ProDom; PD004723; PEPCK_ATP; 1.
 DR TIGRFAMS; TIGR00224; PCKA; 1.
 DR PROSITE; PS00532; PEPCK_ATP; 1.
 KW Gluconeogenesis; Lyase; Decarboxylase; ATP-binding; Complete proteome.

FT NP_BIND 227 234 ATP (BY SIMILARITY).
 SQ SEQUENCE 521 AA; 58771 MW; 1783A3320B106341 CRC64;

Query Match 35.6%; Score 50.5; DB 1; Length 521;
 Best Local Similarity 30.8%; Pred. No. 9.9;
 Matches 8; Conservative 9; Mismatches 8; Indels 1; Gaps 1;

QY 1 NLWA-AQEGRELRMSDEVDSEFKK 25
 |:|:|: |:|:|: |:|:|:|
 Db 481 NTWKDEYDKTAKLAQRFIENFQK 506

RESULT 7

AROG_YEAST
 ID AROG_YEAST STANDARD; PRT; 370 AA.
 AC P32449;
 DT 01-OCT-1993 (Rel. 27, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Phospho-2-dehydro-3-deoxyheptonsate aldolase, tyrosine-inhibited
 DE (EC 4.1.2.15) (Phospho-2-keto-3-deoxyheptonsate aldolase) (DHAP
 DE synthetase) (3-deoxy-D-arabino-heptulosonate 7-phosphate synthase).
 GN ARO4 OR YBR249C OR YBR1701.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=9225349; PubMed=1348717;
 RX Kuenzler M., Paravicini G., Egli C., Irniger S., Braus G.H.;
 RT "Cloning, primary structure and regulation of the ARO4 gene, encoding
 RT the tyrosine-inhibited 3-deoxy-D-arabino-heptulosonate-7-phosphate
 RT synthase from Saccharomyces cerevisiae.";
 RL Gene 113:67-74(1992).
 RN [2]
 RP REVISIONS TO 205-207.
 RA Kuenzler M.;
 RL Submitted (NOV-1993) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RX MEDLINE=94078675; PubMed=8256522;
 RA Daignon F., Biteau N., Aigle M., Crouzet M.;
 RT "The complete sequence of a 6794 bp segment located on the right arm
 RT of chromosome II of Saccharomyces cerevisiae. Finding of a putative
 RT dUTPase in a yeast.";
 RL Yeast 9:1131-1137(1993).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RA Aljinovic G., Pohl F.M., Pohl T.M.;
 RL Submitted (AUG-1994) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: STEREOSPECIFIC CONDENSATION OF PHOSPHENOLPYRUVATE (PEP)
 CC AND D-ERYTHROSE-4-PHOSPHATE (E4P) GIVING RISE TO 3-DEOXY-D-
 CC ARABINO-HEPTULOSONATE-7-PHOSPHATE (DAHP).
 CC -!- CATALYTIC ACTIVITY: 2-dehydro-3-deoxy-D-arabino-heptonsate 7-
 CC phosphate + phosphate = phosphoenolpyruvate + D-erythrose 4-
 CC phosphate + H(2O).
 CC -!- ENZYME REGULATION: INHIBITED BY TYROSINE.
 CC -!- PATHWAY: Aromatic amino acids biosynthesis; shikimate pathway;
 CC first step.
 CC -!- INDUCTION: By amino acid starvation.
 CC -!- SIMILARITY: BELONGS TO CLASS-I DAHP SYNTHETASE FAMILY.
 CC
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DR EMBL; X61107; CAA43419.1; -.
DR EMBL; L20296; CAA65607.1; -.
DR EMBL; Z36118; CAA85212.1; -.
DR PIR; S38185; S38185.
DR HSP; P00886; IQR7.
DR SGD; S0000453; ARO4.
DR GO; GO:0003849; F:2-dehydro-3-deoxyphosphoheptonate aldolase . . .; IDA.
DR InterPro; IPR006219; AROFGH.
DR InterPro; IPR006218; DAHP1/KDSA.
DR Pfam; PF00793; DAHP-synth_1; 1.
DR Pfam; PF005060; AROFGH; 1.
DR TIGRFAMS; TIGR00034; aroFGH; 1.
KW Aromatic amino acid biosynthesis; Lyase; Multigene family.
SQ SEQUENCE 370 AA; 39749 MW; 594ED48F24175979 CRC64;

Query Match 35.28; Score 50; DB 1; Length 370;
Best Local Similarity 55.68; Pred. No. 8.1;
Matches 10; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 1 NLWAAQEGYGLRMSDE 18
      :| ||||| |::|||
Db 80 DLRAQEVALLKLSDE 97

RESULT 8
BXDL_DROME
ID BXDL_DROME STANDARD; PRT; 320 AA.
AC Q9VE53; Q95U84;
DT 16-OCT-2001 (Rel. 40, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Brix domain containing protein 1 homolog.
GN CG7993.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkley;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Ananides P.G., Scher S.E., Li P.W., Hoskins R.A., Galie R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.D.G.,
RA April J.F., Adayani A., An H.-J., Andrews-Fianknoch C., Baldwin D.,
RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,
RA Burtis K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferrieria S., Fleischmann W.,
RA Fogle K., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Helman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McInosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milshina N.V., McCarthy C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Sigen-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,

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RA Wang Z.-Y., Wassarman D.A., Weinstock G.M., Weissenbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkley;
RX MEDLINE=22426066; PubMed=12537569;
RA Stapleton M., Carlson J.W., Brokstein P., Yu C., Champe M.,
RA George R.A., Guarin H., Kronmiller B., Pacleb J.M., Park S., Wan K.H.,
RA Rubin G.M., Celniker S.E.;
RT "A Drosophila full-length cDNA resource.";
RL Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8(2002).
CC -!- SUBCELLULAR LOCATION: Nuclear; nucleolar (By similarity).
CC -!- SIMILARITY: Contains 1 Brix domain.
CC -!- CAUTION: Ref.1 sequence differs from that shown due to erroneous
CC gene model prediction.
CC -----
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CC -----
DR EMBL; AE003721; AAF5514.1; ALT_SEQ.
DR EMBL; AY058248; AAL13477.1; -.
DR FlyBase; FBgn0038585; CG7993.
DR InterPro; IPR007109; Brix.
DR Pfam; PF04427; Brix; 1.
KW Hypothetical protein; Nuclear protein.
FT DOMAIN 30 243 BRIX.
FT CONFLICT 188 188 I -> T (IN REF. 1).
SQ SEQUENCE 320 AA; 36509 MW; EE98936DD68B3703 CRC64;

Query Match 34.98; Score 49.5; DB 1; Length 320;
Best Local Similarity 43.58; Pred. No. 8.1;
Matches 10; Conservative 6; Mismatches 4; Indels 3; Gaps 1;

QY 3 WAAQEGYGLRMSDEFVDFKK 25
      || ||||| :|::|:
Db 149 WQATE---ELRLRLMLFDTFQR 168

RESULT 9
HXK_DEBOC
ID HXK_DEBOC STANDARD; PRT; 478 AA.
AC P50506;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE Hexokinase (EC 2.7.1.1).
GN HXK.
OS Debaryomyces occidentalis (Yeast) (Schwanniomyces occidentalis).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Debaryomyces.
OX NCBI_TaxID=27300;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=APCC 2322 / CBS 819;
RX MEDLINE=95339407; PubMed=7614556;
RA Rose M.;
RT "Molecular and biochemical characterization of the hexokinase from
RT the starch-utilizing yeast Schwanniomyces occidentalis.";
RL Curr. Genet. 27:330-338(1995).
CC -!- CATABOLIC ACTIVITY: ATP + D-hexose = ADP + D-hexose 6-phosphate.
CC -!- PATHWAY: FIRST STEP OF SEVERAL METABOLIC PATHWAYS.
CC -!- SUBUNIT: Homodimer (By similarity).
CC -!- SIMILARITY: BELONGS TO THE HEXOKINASE FAMILY.

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CC -----
CC EMBL: S78714; AAB34892.1; -.
CC PIR: S57203; S57203.
CC HSSP: Q26609; IBDG.
CC InterPro: IPR001312; Hexokinase.
CC Pfam: PF03737; hexokinase2; 1.
CC Pfam: PF00349; hexokinase; 1.
CC PRINTS: PR00475; HEXOKINASE.
CC PRODOM: PD001109; Hexokinase; 1.
CC PROSITE: PS00378; HEXOKINASES; 1.
CC TRANSFERASE: Kinase; Glycolysis; Allosteric enzyme; ATP-binding.
CC BINDING 111 111 ATP (BY SIMILARITY)
CC DOMAIN 151 177 GLUCOSE-BINDING (POTENTIAL).
CC SEQUENCE 478 AA; 53066 MW; 080D5F9134478ABA CRC64;
CC -----
Query Match 34.5%; Score 49; DB 1; Length 478;
Best Local Similarity 42.1%; Pred. No. 15;
Matches 8; Conservative 5; Mismatches 6; Indels 0; Gaps 0;
QY 9 GRELRMSDFVDSFKGL 27
DB 38 GETLRITDFEISELEKGL 56
CC -----
RESULT 10
AROG_CANAL STANDARD; PRT; 370 AA.
AC P79023;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Phospho-2-dehydro-3-deoxyheptanate aldolase, tyrosine-inhibited
DE (EC 4.1.2.15) (Phospho-2-keto-3-deoxyheptanate aldolase) (DAHP
DE synthetase) (3-deoxy-D-arabino-heptulosonate 7-phosphate synthase).
GN AR04.
OS Candida albicans (Yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; mitosporic Saccharomycetales; Candida.
OX NCBI_TaxID=3476;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 11651 / B792;
RA Sousa S., Pereira S.A., Livi G.P.;
RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP PARTIAL SEQUENCE FROM N.A.
RX MEDLINE=96207468; PubMed=8625423;
RA Pereira S.A., Livi G.P.;
RL "Aromatic amino-acid biosynthesis in Candida albicans: identification
RT of the AR04 gene encoding a second DAHP synthase.";
RL Curr. Genet. 29:441-445(1996).
CC -!- FUNCTION: STEREOSPECIFIC CONDENSATION OF PHOSPHOENOLPYRUVATE (PEP)
CC AND D-ERYTHROSE-4-PHOSPHATE (E4P) GIVING RISE TO 3-DEOXY-D-
CC ARABINO-HEPTULOSONATE-7-PHOSPHATE (DAHP).
CC -!- CATALYTIC ACTIVITY: 2-dehydro-3-deoxy-D-arabino-heptanate 7-
CC phosphate + phosphate = phosphoenolpyruvate + D-erythrose 4-
CC phosphate + H(2)O.
CC -!- ENZYME REGULATION: INHIBITED BY TYROSINE (BY SIMILARITY).
CC -!- PATHWAY: Aromatic amino acids biosynthesis; shikimate pathway;
CC first step.
CC -!- SIMILARITY: BELONGS TO CLASS-I DAHP SYNTHETASE FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: U53216; AAB48240.1; -.
CC HSSP: P00886; IQR7.
CC InterPro: IPR006219; AROFGH.
CC InterPro: IPR006218; DAHP1/ADSA.
CC Pfam: PF00793; DAHP_synth_1; 1.
CC PRODOM: PD005060; AROFGH; 1.
CC TIGRFAMs: TIGR00034; aroFGH; 1.
CC Aromatic amino acid biosynthesis; Lysine; Multigene family.
CC SEQUENCE 370 AA; 40291 MW; 11E5B324C8D7B6DB CRC64;
CC -----
Query Match 33.8%; Score 48; DB 1; Length 370;
Best Local Similarity 47.1%; Pred. No. 16;
Matches 8; Conservative 5; Mismatches 4; Indels 0; Gaps 0;
QY 5 AQEYGRELRMSDFVD 21
DB 74 ALEYGRLKXLDELKD 90
CC -----
RESULT 11
VP5_BTIV2A STANDARD; PRT; 526 AA.
AC P30209;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Outer capsid protein VP5.
GN S6.
OS Bluetongue virus (serotype 2 / isolate USA).
OC Viruses; dsRNA viruses; Reoviridae; Orbivirus.
OX NCBI_TaxID=10907;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90209359; PubMed=2157314;
RA Hirasawa T., Roy P.;
RL "The complete nucleotide sequence of VP5 of a strain of bluetongue
RT virus of serotype 2 isolated in the USA reveals its close
RT relationship with a virus of serotype 1 isolated in Australia.";
RL Virus Res. 15:107-112(1990).
CC -!- FUNCTION: THE VP5 PROTEIN IS ONE OF THE TWO PROTEINS (WITH VP2)
CC WHICH CONSTITUTE THE VIRUS PARTICLE OUTER CAPSID.
CC -!- SIMILARITY: BELONGS TO THE REOVIRUSES VP5 FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: X62283; CAA44172.1; -.
CC PIR: A43486; P5XRBU.
CC InterPro: IPR000145; Orbi_VP5.
CC Pfam: PF00901; Orbi_VP5; 1.
CC Coat protein.
CC SEQUENCE 526 AA; 58953 MW; DE50D6013B983A04 CRC64;
CC -----
Query Match 33.8%; Score 48; DB 1; Length 526;
Best Local Similarity 41.7%; Pred. No. 23;
Matches 10; Conservative 4; Mismatches 10; Indels 0; Gaps 0;
QY 4 AAQEGYGRELRMSDFVDSFKKGL 27
DB 309 AIQENHKELMHINDEILPRFKAM 332
CC -----
RESULT 12

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KW	RNA-binding; tRNA-binding; Metal-binding; Zinc; Complete proteome.
FT SITE	10 20 "HIGH" REGION. FT SITE 297 301 "KMSKS" REGION.
FT DOMAIN	529 629 TRNA-BINDING
FT METAL	125 125 ZINC (BY SIMILARITY).
FT METAL	128 128 ZINC (BY SIMILARITY).
FT METAL	146 146 ZINC (BY SIMILARITY).
FT METAL	149 149 ZINC (BY SIMILARITY).
FT BINDING	300 300 ATP (BY SIMILARITY).
SQ SEQUENCE	629 AA; 73004 MW; BOE0759F7C78ACEC CRC64;
Query Match	33.8%; Score 48; DB 1; Length 629;
Best Local Similarity	37.5%; Pred.No. 28;
Matches	9; Conservative 6; Mismatches 9; Indels 0; Gaps 0;
QY	2 LWRRAOEGRELRMSDFVDSFKK 25 : :: : : :
Dd	55 LQAAGAKGDPQEFCDLAERPKR 78 : :: : : :
RESULT 13	
PPCK_RHISN	STANDARD; PRT; 537 AA.
ID PPCK_RHISN	
AC F43086;	
DT 01-NOV-1995 (Rel. 32, Created)	
DT 01-NOV-1995 (Rel. 32, Last sequence update)	
DT 28-FEB-2003 (Rel. 41, Last annotation update)	
DE Phosphoenolpyruvate carboxykinase [ATP] (EC 4.1.1.49) (PEP	
DE carboxykinase) (Phosphoenolpyruvate carboxylase) (PFCK).	
GN PCKA.	
OS Rhizobium sp. (strain NGR234).	
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;	
NC Ribobiaceae; Rhizobium/agrobacterium group; Rhizobium.	
OX NCBI_TaxID=394;	
XN [1]	
RP SEQUENCE FROM N.A.	
RX MEDLINE=92079005; PubMed=1720862;	
RA Oesterhaas M., Finan T.M., Stanley J.;	
RT "Site-directed mutagenesis and DNA sequence of pcka of Rhizobium	
NCR234, encoding phosphoenolpyruvate carboxykinase; gluconeogenesis	
RI and host-dependent symbiotic phenotype." ;	
RL Mol. Gen. Genet. 230:257-269(1991).	
CC -I CATALYTIC ACTIVITY: ATP + oxaloacetate = ADP + phosphoenolpyruvate	
CC + CO(2).	
CC -I PATHWAY: Rate-limiting gluconeogenic enzyme.	
CC -I SUBCELLULAR LOCATION: Cytoplasmic (By similarity).	
CC -I SIMILARITY: Belongs to the phosphoenolpyruvate carboxykinase [ATP]	
CC family.	
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DR ENBL; X63291; CAA44925.1; -	
DR HSSP; P22359; lAQ2.	
DR HAMAP; MF_00453; -. 1.	
DR InterPro; IPR001272; PEPC_K_ATP.	
DR Pfam; PF01293; PEPC_K_ATP; 1.	
DR ProDom; PD004723; PEPC_K_ATP; 1.	
DR TIGRFAMs; TIGR00224; pckA; 1.	
DR PROSITE; PS00532; PEPC_K_ATP; 1.	
KW Glucoconegensis; Lyase; Decarboxylase; ATP-binding.	
FT NP_BIND 236 243 ATP (BY SIMILARITY).	
SQ SEQUENCE 537 AA; 58370 MW; B2CED7FA54326B1E CRC64;	
Query Match	33.5%; Score 47.5; DB 1; Length 537;
Best Local Similarity	41.7%; Pred. No. 28;
Matches	10; Conservative 5; Mismatches 8; Indels 1; Gaps 1;


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QY 3 WAA-QYEGRELRRMSDFVDSFKK 25
   ||||| : ||| : ||| : |||
Db 493 WSRGQAYDAQARLYDMFIANFAK 516

RESULT 14
VG13_BPMD2
ID VG13_BPMD2 STANDARD; PRT; 595 AA.
AC 064206;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Gene 13 protein (GP13).
GN 13.
OS Mycobacteriophage D29.
OC Viruses; dsDNA viruses, no RNA stage; Caudovirales; Siphoviridae.
OX NCBI_TaxID=28369;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98300335; PubMed=9636706;
RA Ford M.E., Sarkis G.J., Belanger A.E., Hendrix R.W., Hatfull G.F.;
RT "Genome structure of mycobacteriophage D29: implications for phage
   evolution.";
RL J. Mol. Biol. 279:143-164(1998).
CC -!- SIMILARITY: BELONGS TO THE PHAGE TERMINASE FAMILY.
CC
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   or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF022214; AAC18453.1; -.
DR PIR; B72801; B72801.
DR InterPro; IPR005021; Phage_term1n.
DR Pfam; PF03354; Phage_terminase; 1.
SQ SEQUENCE 595 AA; 65397 MW; AFD123ED5371E263 CRC64;

Query Match 33.5%; Score 47.5; DB 1; Length 595;
Best Local Similarity 45.5%; Pred. No. 31;
Matches 10; Conservative 5; Mismatches 4; Indels 3; Gaps 1;

QY 1 NLWAAQYEGRELRRMSDFVDS 22
   ::||| : ||| : ||| :
Db 435 DIWDPOKYGGEVFR---EFVDA 453

RESULT 15
RPSD_BORBU
ID RPSD_BORBU STANDARD; PRT; 631 AA.
AC P52323;
DT 01-OCT-1996 (Rel. 34, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE RNA polymerase sigma factor rpoD (Sigma-70).
GN RPOD OR BE0712.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE OF 89-631 FROM N.A.
RC STRAIN=ATCC 35210 / B31;
RA Pan M.;
RL Thesis (1994), National Taiwan University, Taiwan.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 35210 / B31;
RX MEDLINE=98065943; PubMed=9403685;
RA Fraser C.M., Casjens S., Huang W.M., Sutton G.G., Clayton R.A.,
RA Lathigra R., White O., Ketchum K.A., Dodson R., Hickey E.K., Gwinn M.,
RA Dougherty B., Tomb J.-F., Fleischmann R.D., Richardson D.,

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RA Peterson J., Kervage A.R., Quackenbush J., Salzberg S., Hanson M.,
RA van Vugt R., Palmer N., Adams M.D., Gocayne J.D., Weidman J.,
RA Utterback T., Watthey L., McDonald L., Artiach P., Bowman C.,
RA Garland S., Fujii C., Cotton M.D., Horst K., Roberts K., Hatch B.,
RA Smith H.O., Venter J.C.;
RT "Genomic sequence of a Lyme disease spirochaete, Borrelia
   burgdorferi.";
RL Nature 390:580-586(1997).
RN [3]
RP SEQUENCE OF 165-614 FROM N.A.
RC STRAIN=297;
RA Pan M., Yeh J., Tsai C.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ Databases.
CC -!- FUNCTION: THE SIGMA FACTOR IS AN INITIATION FACTOR THAT PROMOTES
   ATTACHMENT OF THE RNA POLYMERASE TO SPECIFIC INITIATION SITES AND
   THEN IS RELEASED. THIS IS THE PRIMARY SIGMA-FACTOR OF THIS
   BACTERIA.
CC
CC -!- SIMILARITY: Belongs to the sigma-70 factor family.
CC
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CC -----
CC EMBL; U17591; AAC44104.1; -.
DR PIR; AE001171; AAC67061.1; -.
DR EMBL; U68006; AAC45100.1; -.
DR PIR; G70188; G70188.
DR HSP; P00579; ISIG.
DR TIGR; BB0712; -.
DR InterPro; IPR000943; Sigma_70.
DR Pfam; PF03979; sigma70_x1_1; 1.
DR Pfam; PF00140; sigma70_x1_2; 1.
DR Pfam; PF04542; sigma70_x2; 1.
DR Pfam; PF04539; sigma70_x3; 1.
DR Pfam; PF04545; sigma70_x4; 1.
DR PROSITE; PS00715; SIGMA70_1; 1.
DR PROSITE; PS00716; SIGMA70_2; 1.
DR PROSITE; PS00716; SIGMA70_2; 1.
KW Transcription regulation; Sigma factor; DNA-directed RNA polymerase;
FT DOMAIN 419 437 POLYMERASE CORE BINDING (POTENTIAL).
FT DNA_BIND 589 608 H-T-H MOTIF (BY SIMILARITY).
SQ SEQUENCE 631 AA; 73642 MW; BD565AB7D8F44796 CRC64;

Query Match 33.1%; Score 47; DB 1; Length 631;
Best Local Similarity 50.0%; Pred. No. 39;
Matches 9; Conservative 3; Mismatches 6; Indels 0; Gaps 0;

QY 8 YGRELRRMSDFVDSFKK 25
   | ||| ||| : ||| :
Db 279 YQEELRIFSDDYIDSANK 296

Search completed: September 23, 2003, 09:47:46
Job time : 26 secs

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GenCore version 5.1.1.6
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OM protein - protein search, using sw model

Run on: September 23, 2003, 09:43:16 : Search time 96 Seconds
(without alignments)
72.577 Million cell updates/sec

Title: US-09-544-664b-2

Perfect score: 142

Sequence: 1 NLWAAQEGRELRMSDEFVDSFKKL 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_23:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_virus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	84	59.2	146	13 Q9I9N2	Q9I9N2 brachydanio
2	54	38.0	606	10 Q8H4M3	Q8H4M3 oryza sativ
3	53	37.3	85	16 Q8R8Y1	Q8R8Y1 thermoanaer
4	52	36.6	889	10 Q8LMF2	Q8LMF2 oryza sativ
5	51	35.9	335	10 Q64692	Q64692 arabidopsis
6	51	35.9	722	17 Q8TVX8	Q8TVX8 methanopyru
7	51	35.9	1248	16 Q9HZQ3	Q9HZQ3 pseudomonas
8	50	35.2	173	4 Q8N955	Q8N955 homo sapien
9	50	35.2	230	16 Q8XXS6	Q8XXS6 raistonia s
10	50	35.2	330	16 Q8FSR1	Q8FSR1 leptospira
11	50	35.2	363	16 Q8EH69	Q8EH69 shewarella
12	50	35.2	371	5 Q8U1L1	Q8U1L1 drosophila
13	50	35.2	369	5 Q9WSB6	Q9WSB6 drosophila
14	49.5	34.9	320	5 Q9SU84	Q9SU84 drosophila
15	49	34.5	214	16 Q8KDC3	Q8KDC3 chlorobium
16	49	34.5	247	2 Q9AJL5	Q9AJL5 streptomyce

17	49	34.5	378	17	Q8PYK0	Q8PYK0 methanosarc
18	49	34.5	564	16	Q9RUK9	Q9RUK9 deinococcus
19	49	34.5	1039	17	Q8TME7	Q8TME7 methanosarc
20	49	34.5	1350	5	O17342	O17342 caenorhabdi
21	49	34.5	1690	17	Q9HKL0	Q9HKL0 thermoplasma
22	49	34.5	1713	17	Q97B15	Q97B15 thermoplasma
23	48.5	34.2	130	16	Q8R937	Q8R937 thermoanaer
24	48.5	34.2	937	16	Q97H37	Q97H37 clostridium
25	48.5	34.2	1303	12	Q9QSU0	Q9QSU0 potato mop-
26	48	33.8	196	16	Q8VUS3	Q8VUS3 mycobacteri
27	48	33.8	223	16	Q10843	Q10843 mycobacteri
28	48	33.8	288	10	Q94ID5	Q94ID5 oryza sativ
29	48	33.8	295	16	Q9HV62	Q9HV62 pseudomonas
30	48	33.8	339	10	Q9F005	Q9F005 atrichum an
31	48	33.8	357	16	Q8PEY8	Q8PEY8 xanthomonas
32	48	33.8	370	10	Q94ID4	Q94ID4 oryza sativ
33	48	33.8	373	10	Q9FU53	Q9FU53 oryza sativ
34	48	33.8	373	10	Q94IE4	Q94IE4 oryza sativ
35	48	33.8	526	12	Q8BAD7	Q8BAD7 bluetongue
36	48	33.8	526	12	Q8BAD6	Q8BAD6 bluetongue
37	48	33.8	709	10	Q8GUT8	Q8GUT8 arabidopsis
38	47.5	33.5	233	17	Q29007	Q29007 archaeoglob
39	47.5	33.5	234	16	Q8ZAP2	Q8ZAP2 yersinia pe
40	47.5	33.5	238	16	Q99U03	Q99U03 staphylococ
41	47.5	33.5	279	10	Q8W4Z1	Q8W4Z1 brassica ol
42	47.5	33.5	309	10	Q9SXJ6	Q9SXJ6 arabidopsis
43	47.5	33.5	309	10	Q94JY6	Q94JY6 arabidopsis
44	47.5	33.5	310	10	Q824Z1	Q824Z1 arabidopsis
45	47.5	33.5	311	10	Q9STI9	Q9STI9 brassica ca

ALIGNMENTS

RESULT 1
Q9I9N2 ID Q9I9N2 PRELIMINARY; PRT; 146 AA.
AC Q9I9N2;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Bad.
GN BAD.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE-20373792; PubMed-10917738;
RA Inohara N., Nunez G.;
RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish."
RL Cell Death Differ. 7:509-510(2000).
DR EMBL; AF231017; AAF66962.2; -
DR HSP; Q92934; IG5J
DR ZFIN; ZDB-GENE-000616-1; bad.
SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;

Query Match 59.2%; Score 84; DB 13; Length 146;

Best Local Similarity 62.5%; Pred. No. 0.00023;

Matches 15; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 2 LWAQEGRELRMSDEFVDSFKK 25

|||||:|||||:|||||:|

Db 89 LWAQYGGQLRMSDEFDKGMR 112

RESULT 2

Q8H4M3

ID Q8H4M3 PRELIMINARY; PRT; 606 AA.

AC Q8H4M3;

OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoidae; Oryzaceae; Oryza.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA McCombie W.R., de la Bastide M., Spiegel L., Preston R., Ferraro K.,
RA Kuit K., Nascimento L., Zutavern T., Ballia V., Bell M., Baker J.,
RA Santos L., Miller B., Katzenberger F., Muller S., King L., Yang C.,
RA Dike S., O'Shaughnessy A., Palmer L., Dedhia N.;
RT "genomic sequence for Oryza sativa, Nipponbare strain, clone
OT CJ1325D05, from chromosome 10, complete sequence.";
RL Submitted (JUL-2002) to the EMBL/GenBank/DDBJ databases.
DR EMBL; AC115686; AA074351.1; -
DR Gramineae; Q8LMF2; -
RW Hypothetical protein.
SQ SEQUENCE 889 AA; 101583 MW; C47D8715883D6376 CRC64;

Query Match 36.6%; Score 52; DB 10; Length 889;
Best Local Similarity 40.0%; Pred.No.79;
Matches 10; Conservative 6; Mismatches 9; Indels 0; Gaps 0;

QY 3 WAAQEXGRELRRMSDEFVDSFKKGL 27
II I :::: :::
DB 627 WALDEAPWEMKRLHTWTYWDASKKGL 651

RESULT 5
ID 064692 PRELIMINARY; PRT; 335 AA.
IC 064692;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Putative GA4 protein.
GN T31E10.11 OR GA20X3.
GC Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsiis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv Columbia;
RA Rounsley S.D., Kaul S., Lin X., Ketchum K.A., Crosby M.L.,
RA Brandon R.C., Sykes S.M., Mason T.M., Kerlavage A.R., Adams M.D.,
RA Scmerville C.R., Venter J.C.;
RL Submitted (MAY-1998) to the EMBL/GenBank/DDBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=92218343; PubMed=10200325;
RA Thomas S.G., Phillips A.L., Hedden P.;
RT "Molecular cloning and functional expression of gibberellin 2-
RT oxidases, multifunctional enzymes involved in gibberellin
RT deactivation";
RL Proc. Natl. Acad. Sci. U.S.A. 96:4698-4703(1999).
DR EMBL; AC004077; AAA14908.1; -
DR EMBL; AJ132437; CAB41009.1; -
DR InterPro; IPR005123; ZOG-FcEl_Oxy.
DR Pfam; PF03171; ZOG-FcEl_Oxy; 1.
SQ SEQUENCE 335 AA; 38216 MW; 181F6EAA1BE1C331 CRC64;

Query Match 35.9%; Score 51; DB 10; Length 335;
Best Local Similarity 37.5%; Pred.No.37;
Matches 9; Conservative .9; Mismatches 6; Indels 0; Gaps 0;

QY 4 AAQYEGRELRRMSDEFVDSFKKGL 27
I :|| :||| :|| :||
DB 136 AVETIKEMKRSSKFLEMWVEEL 159

Matches	11;	Conservative		5; Mismatches		5; Indels		2; Gaps		1;
QY	2	LWAAQEYG--RELRRMSDEFVDS 22 : : :								
DB	611	LTRASYGPLRLERLAFDYDA 633								
RESULT 8										
ID Q8N955	PRELIMINARY; PRT; 173 AA.									
AC Q8N955:										
DT 01-OCT-2002 (TrEMBLrel. 22, Created)										
DT 01-MAR-2002 (TrEMBLrel. 22, Last sequence update)										
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)										
DE Hypothetical protein FLJ38343.										
OS Homo sapiens (human).										
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;										
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.										
NCBI_TaxID=9606;										
RN [1]										
RS SEQUENCE FROM N.A.										
RC TISSUE=Brain;										
RA Kanehori K., Ishibashi T., Chiba Y., Fujimori K., Hiraoka S., Tanai H., Watanabe S., Ishida S., Ono Y., Kotura T., Watanabe M., Sugiyama T., Irie R., Otsuki T., Sato H., Ota F., Wakamatsu A., Ishii S., Yamamoto J., Isono Y., Kawai-Hio Y., Saito K., Nishikawa T., R Kimura K., Matsuo K., Nakamura Y., Sekine M., Kikuchi H., Kanda K., RA Wagatsuma M., Takahashi-Fujii A., Oshima A., Sugiyama A., Kawakami B., RA Suzuki Y., Sugano S., Nagahari K., Masuho Y., Nagai K., Isogai T.; RT "NEO human cDNA sequencing project"; Submitted (JUL-2002) to the EMBL/GenBank/DDBJ databases. NL DBML; AK095662; BAC04601.1; -; KW Hypothetical protein. SQ SEQUENCE 173 AA; 21285 MW; 0EBFF3BFD96FE775 CRC64;										
Query Match	35.2%;	Score 50;	DB 4;	Length 173;						
Best Local Similarity	47.6%;	Pred. No. 24;								
Matches	10;	Conservative	4; Mismatches	7; Indels	0; Gaps					
QY	6	QEYGRLELRMSDFVDSPKKG 26 : ::								
DB	36	KEGGERRKERFWIDGRKKG 56								
RESULT 9										
Q8XXS6	PRELIMINARY; PRT; 230 AA.									
ID Q8XXS6:										
AC Q8XXS6:										
DT 01-WAR-2002 (TrEMBLrel. 20, Created)										
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)										
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)										
DE Probable ATP-binding ABC transporter protein.										
GN RSC2037 OR RS03602.										
OS Ralstonia solanacearum (Pseudomonas solanacearum).										
OC Bacteria; Proteobacteria; betaproteobacteria; Burkholderiales;										
OC Ralstoniaceae; Ralstonia.										
NCBI_TaxID=305;										
RN [1]										
RS SEQUENCE FROM N.A.										
RC STRAIN=GMI1000;										
RX MEDLINE=21681879; PubMed=11823852;										
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S., RA Ariat M., Billault A., Brottier P., Camus J.C., Cattolico L., RA Chandler M., Choise N., Claudel-Renard C., Cunnac S., Demange N., RA Gaspin C., Lavie M., Moisan A., Robert C., Sourin W., Schlex T., RA Siguer P., Thebaud P., Whalen M., Wincker P., Levy M., RA Weissenbach J., Boucher C.A.; RT "Genome sequence of the plant pathogen Ralstonia solanacearum"; FL Nature 415:457-502(2002). DR EMBL; AL646067; CAD15739.1; -; DR InterPro; IPR003593; AAA_AtPase. DR InterPro; IPR003439; ABC_transporter. DR Pfam; PF00005; ABC_tran; 1.										

RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agayani A., An H.-J., Andrews-Prannkoc C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Fostler C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Helman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskas R., Tector K., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.,
 RA "The genome sequence of *Drosophila melanogaster*.",
 RT Science 287:2185-2195(2000).
 RL [2]
 RN
 RP SEQUENCE FROM N.A.
 RA Celniker S.E., Adams M.D., Kronmiller B., Wan K.H., Holt R.A.,
 RA Evans C.A., Gocayne J.D., Anantides P.G., Brandon R.C., Rogers Y.,
 RA Banson J., An H., Baldwin D., Banton J., Beeson K.Y., Busam D.A.,
 RA Carlson J.W., Center A., Champe M., Davenport L.B., Dietz S.M.,
 RA Dodson K., Dorsett V., Doup L.E., Doyle C., Dresnek D., Farfan D.,
 RA Ferreira S., Frise E., Galle R.F., Garg N.S., George R.A.,
 RA Gonzalez M., Houck J., Hoskins R.A., Hostin D., Howland T.J.,
 RA Ibegwam C., Jalali M., Kruse D., Li P., Mattei B., Moshrefi A.,
 RA McIntosh T.C., Moy M., Murphy B., Nelson C., Nelson K.A., Nunoo J.,
 RA Pacle J., Paragas V., Park S., Patel S., Pfeiffer B.,
 RA Phouanavong S., Pittman G.S., Puri V., Richards S., Scheeler F.,
 RA Stapleton M., Strong R., Svirskas R., Tector C., Tyler D.,
 RA Williams S.M., Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.,
 RT "Sequencing of *Drosophila melanogaster* genome".
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Misra S., Crosby M.A., Matthews B.B., Bayraktaroglu L., Campbell K.,
 RA Hradecky P., Huang Y., Kaminker J.S., Prochnik S.E., Smith C.D.,
 RA Tupy J.L., Bergman C., Berman B., Carlson J.W., Celniker S.E.,
 RA Clamp M., Drysdale R., Emmert D., Frise E., de Grey A., Harris N.,
 RA Kronmiller B., Marshall B., Millburn G., Richter J., Russo S.,
 RA Searle S.M.J., Smith E., Shu S., Smutniak F., Whitfield E.,
 RA Ashburner M., Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.,
 RT "Annotation of *Drosophila melanogaster* genome".
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RA Adams M.D., Celniker S.E., Gibbs R.A., Rubin G.M., Venter C.J.,
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE FROM N.A.
 RA FlyBase.
 RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RP SEQUENCE FROM N.A.

RC STRAIN=Berkeley;
 RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
 RA Champe M., Chavez C., Dorsett V., Farfan D., Frise E., George R.,
 RA Gonzalez M., Guarin H., Li P., Liao G., Miranda A., Mungall C.J.,
 RA Nunoo J., Pacle J., Paragas V., Park S., Phouanavong S., Wan K.,
 RA Yu C., Lewis S.E., Rubin G.M., Celniker S.,
 RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AE003419; AAF45578.3; -;
 DR EMBL; AY051773; AAK93197.1; -;
 DR FlyBase; FBgn0040337; EG:BACR7A4.8.
 DR InterPro; IPR004506; TrmU.
 DR Pfam; PF03054; tRNA_Me_trans; 1.
 DR TIGRFAMs; TIGR00420; trmU; 1.
 SQ SEQUENCE 389 AA; 43350 MW; 20C9405FFCC9FAE2 CRC64;
 Query Match 35.2%; Score 50; DB 5; Length 389;
 Best Local Similarity 28.6%; Pred. No. 61;
 Matches 10; Conservative 9; Mismatches 6; Indels 10; Gaps 1;
 QY 3 WAAQCYGRELRMS-----DEFVDSFKKGL 27
 Db 57 WACQLGVELRQVNVYREYWTAVFSQFLDDYQML 91
 RESULT 14
 Q95U84 PRELIMINARY; PRT; 320 AA.
 AC Q95U84;
 DT 01-DEC-2001 (TrEMBLrel. 19, Created)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE GH01229p.
 GN CG7993
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkeley;
 RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
 RA Champe M., Chavez C., Dorsett V., Farfan D., Frise E., George R.,
 RA Gonzalez M., Guarin H., Li P., Liao G., Miranda A., Mungall C.J.,
 RA Nunoo J., Pacle J., Paragas V., Park S., Phouanavong S., Wan K.,
 RA Yu C., Lewis S.E., Rubin G.M., Celniker S.,
 RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY058248; AAL13477.1; -;
 DR FlyBase; FBgn0038585; CG7993.
 DR InterPro; IPR007109; Brix.
 DR Pfam; PF04427; Brix; 1.
 DR PROSITE; PS0833; BRIX; 1.
 SQ SEQUENCE 320 AA; 36509 MW; EE98936DD68B3703 CRC64;
 Query Match 34.9%; Score 49.5; DB 5; Length 320;
 Best Local Similarity 43.5%; Pred. No. 58;
 Matches 10; Conservative 6; Mismatches 4; Indels 3; Gaps 1;
 QY 3 WAAQCYGRELRMSDEFVDSFKK 25
 Db 149 WAQTE--ELRLRLNLFIDTQR 168
 RESULT 15
 Q8KDC3 PRELIMINARY; PRT; 214 AA.
 ID Q8KDC3;
 AC Q8KDC3;
 DT 01-OCT-2002 (TrEMBLrel. 22, Created)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Hypothetical protein CTL131.
 GN CTL131.
 OS Chlorobium tepidum.

```

OC Bacteria; Chlorobi; Chlorobia; Chlorobiales; Chlorobiaceae;
OC Chlorobium.
OX NCBL_TaxID=1097;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=TLS / ATCC 49652 / DSM 12025;
RX MEDLINE=22103685; PubMed=12093901;
RA Eisen J.A., Nelson K.E., Paulsen I.T., Heidelberg J.F., Wu M.,
RA Dodson R.J., Deboy R., Gwinn M.L., Nelson W.C., Haft D.H.,
RA Hickey E.K., Peterson J.D., Durkin A.S., Kolonay J.L., Yang F.,
RA Holt I., Umayam L.A., Mason T., Brenner M., Shea T.P., Parksey D.,
RA Niernan W.C., Feldblyum T.V., Hansen C.L., Craven M.B., Radune D.,
RA Vanathavan J., Khouri H., White O., Gruber T.M., Ketchum K.A.,
RA Venter J.C., Tettelin H., Bryant D.A., Fraser C.M.; Kirsch J.A.,
RT "The complete genome sequence of Chlorobium tepidum TLS, a
RT photosynthetic, anaerobic, green-sulfur bacterium."
RL Proc. Natl. Acad. Sci. U.S.A. 99:9509-9514(2002).
DR EMBL; AE012874; AAM72364.1; -.
DR TIGR; CT1131; -.
DR InterPro; IPR002785; DUF83.
DR TIGRFAMs; TIGR00372; TIGR00372; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 214 AA; 24004 MW; 56D389219D60B6AD CRC64;

Query Match      34.5%; Score 49; DB 16; Length 214;
Best Local Similarity 37.0%; Pred. No. 43;
Matches 10; Conservative 8; Mismatches 7; Indels 2; Gaps 1;

QY 1 NLWAAQGYGRELRLKMSDFVDSFKKGL 27
   ||: ||: ||: ||: ||: ||: ||:
Db 35 NLYTAE--GREMHERADSAVTSYREGV 59

```

Search completed: September 23, 2003, 09:45:06
 Job time : 101 secs

Result No.	Score	% Match	Query Length	DB ID	Description
1	145	100.0	27	21	Bcl2 polypeptide B
2	145	100.0	162	22	Shorter murine BAD
3	145	100.0	204	17	bcl-x(l)/bcl-2 ass
4	145	100.0	204	17	Murine BCL-XL/BCL-
5	145	100.0	204	19	Mutant BCL-XL/BCL-
6	145	100.0	204	19	Mutant BCL-XL/BCL-
7	145	100.0	204	19	Mutant BCL-XL/BCL-
8	145	100.0	204	19	Murine BAD protein
9	145	100.0	204	22	Longer murine BAD

PD 09-MAY-1996.
 XX
 PF 31-OCT-1995; 95WO-US14246.
 XX
 PR 31-OCT-1994; 94US-0333565.
 XX
 PA (UNIW) UNIV WASHINGTON.
 XX
 PI Korsmeyer SJ;
 XX
 DR WPI; 1996-251465/25.
 DR N-PSDB; AAT29479.
 XX
 PT Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
 PT useful to treat neoplasia and apoptosis and to identify agents
 PT inhibiting its binding to bcl-2 or bcl-x(L) to form heterodimers
 XX
 PS Claim 3; Fig 1; 130pp; English.
 XX
 CC This sequence represents the murine bcl-x(L)/bcl-2 associated death
 CC promoter (Bad) gene. Bad is a 22.1 kD protein which interacts with
 CC bcl-2 and bcl-x proteins and regulates cell death. It has homology
 CC to the bcl-2-related family clustered in the BH1 and BH2 domain. Bad
 CC has been found to hybridise to bcl-x(L) and bcl-2 in yeast two-hybrid
 CC assays and in vivo in mammalian cells. Overexpressed Bad counters the
 CC death inhibitory activity of bcl-x(L), but is much less effective at
 CC countering the death inhibitory activity of bcl-2. Bad expression can
 CC accelerate apoptotic cell death induced by cytokine deprivation in an
 CC IL-3 dependent cell line expressing bcl-x(L), and its also counters the
 CC death repressor activity of bcl-x(L). Bad competes with Bax for binding
 CC to bcl-x(L). Bad may be used to identify agents which inhibit its
 CC binding to bcl-2 or bcl-x(L) to form heterodimers. Such agents may be
 CC used to treat neurodegenerative diseases, immunodeficiency diseases,
 CC e.g. AIDS, senescence or ischaemia.
 XX
 SQ Sequence 204 AA;
 Query Match 100.0%; Score 145; DB 17; Length 204;
 Best Local Similarity 100.0%; Pred. No. 2.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 NLWAAQRYGRELRRMSDEFGSFKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRRMSDEFGSFKGLP 166
 RESULT 4
 AAW61315
 ID AAW61315 standard; Protein: 204 AA.
 XX
 AC AAW61315;
 XX
 DT 07-OCT-1998 (first entry)
 XX
 DE Murine BCL-XL/BCL-2 associated cell death regulator.
 XX
 KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX
 OS Mus sp.
 OS Synthetic.
 XX
 PN WO9817682-A1.
 XX
 PD 30-APR-1998.
 XX
 PF 17-OCT-1997; 97WO-US19175.
 XX
 PR 18-OCT-1996; 96US-0733505.
 XX
 XX (UNIW) UNIV WASHINGTON.
 PA Korsmeyer SJ;
 XX
 PI Korsmeyer SJ;
 XX
 DR WPI; 1998-261422/23.

DR WPI; 1998-261422/23.
 DR N-PSDB; AAV27833.
 XX
 PT New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 XX
 PS Claim 1; Fig 10; 95pp; English.
 XX
 CC The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence is the murine BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX
 SQ Sequence 204 AA;
 Query Match 100.0%; Score 145; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 2.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 NLWAAQRYGRELRRMSDEFGSFKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRRMSDEFGSFKGLP 166
 RESULT 5
 AAW61316
 ID AAW61316 standard; Protein: 204 AA.
 XX
 AC AAW61316;
 XX
 DT 07-OCT-1998 (first entry)
 XX
 DE Mutant BCL-XL/BCL-2 associated cell death regulator #1.
 XX
 KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX
 OS Mus sp.
 OS Synthetic.
 XX
 PN WO9817682-A1.
 XX
 PD 30-APR-1998.
 XX
 PF 17-OCT-1997; 97WO-US19175.
 XX
 PR 18-OCT-1996; 96US-0733505.
 XX
 XX (UNIW) UNIV WASHINGTON.
 PA Korsmeyer SJ;
 XX
 PI Korsmeyer SJ;
 XX
 DR WPI; 1998-261422/23.

DR N-PSDB; AAV27834.

XX New mutant BAD polypeptide with phosphorylatable serine replaced -

PT useful for, e.g. treating reduced apoptosis such as in cancer or

PT viral infection

XX

PS Claim 7; Page 59; 95pp; English.

XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell

CC death regulator) proteins, having an amino acid other than Ser at

CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The

CC present sequence represents a mutant BAD protein. Also described are: (1)

CC fragments of mutant BAD protein able to decrease cell viability; (2)

CC fusion proteins of mutant BAD with a heterologous polypeptide that

CC increases intracellular delivery. Mutant BAD proteins are used to treat

CC or prevent diseases associated with reduced apoptosis, e.g. cancer,

CC viral infection, lymphoproliferation, arthritis, infertility,

CC inflammation and autoimmune disease. Polynucleotide sequences encoding

CC mutant BAD proteins can be used similarly by gene therapy or to produce

CC transgenic animals for use as disease models or in drug screening. BAD

CC proteins phosphorylated at specified Ser are used to screen for enhancers

CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful

CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,

CC aging or ischaemic cell death. The apoptotic status of cells is

CC determined by measuring relative amounts of phosphorylated and non-

CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have

CC greater death-promoting activity than wild-type BAD which can become

CC phosphorylated on the specified Ser, forming a product that does not

CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family

CC proteins in the cytosol, thus promoting cell survival. The mutants with

CC Ser substituted cannot bind 14-3-3.

XX

XX Sequence 204 AA;

SQ

Query Match 100.0%; Score 145; DB 19; Length 204;

Best Local Similarity 100.0%; Pred. No. 2.4e-14;

Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSFKGLP 27

Db 140 NLWAAQRYGRELRLMSDEFGSFKGLP 166

|||||

RESULT 6

AAW61317

ID AAW61317 standard; Protein; 204 AA.

XX

AC AAW61317;

XX

DT 07-OCT-1998 (first entry)

XX

DE Mutant BCL-XL/BCL-2 associated cell death regulator #2.

XX

KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;

KW serine substituted mutant; apoptosis; cancer; viral infection.

XX

OS Mus sp.

OS Synthetic.

XX

PN WO9817682-A1.

XX

PD 30-APR-1998.

XX

PF 17-OCT-1997; 97WO-US19175.

XX

PR 18-OCT-1996; 96US-0733505.

XX

XX (UNIW) UNIV WASHINGTON.

XX

XX Korsmeyer SJ;

XX

XX WPI; 1998-261422/23.

DR N-PSDB; AAV27835.

DR

XX

PT New mutant BAD polypeptide with phosphorylatable serine replaced -

PT useful for, e.g. treating reduced apoptosis such as in cancer or

PT viral infection

XX

PS Claim 7; Page 60; 95pp; English.

XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell

CC death regulator) proteins, having an amino acid other than Ser at

CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The

CC present sequence represents a mutant BAD protein. Also described are: (1)

CC fragments of mutant BAD protein able to decrease cell viability; (2)

CC fusion proteins of mutant BAD with a heterologous polypeptide that

CC increases intracellular delivery. Mutant BAD proteins are used to treat

CC or prevent diseases associated with reduced apoptosis, e.g. cancer,

CC viral infection, lymphoproliferation, arthritis, infertility,

CC inflammation and autoimmune disease. Polynucleotide sequences encoding

CC mutant BAD proteins can be used similarly by gene therapy or to produce

CC transgenic animals for use as disease models or in drug screening. BAD

CC proteins phosphorylated at specified Ser are used to screen for enhancers

CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful

CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,

CC aging or ischaemic cell death. The apoptotic status of cells is

CC determined by measuring relative amounts of phosphorylated and non-

CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have

CC greater death-promoting activity than wild-type BAD which can become

CC phosphorylated on the specified Ser, forming a product that does not

CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family

CC proteins in the cytosol, thus promoting cell survival. The mutants with

CC Ser substituted cannot bind 14-3-3.

XX

XX Sequence 204 AA;

SQ

Query Match 100.0%; Score 145; DB 19; Length 204;

Best Local Similarity 100.0%; Pred. No. 2.4e-14;

Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSFKGLP 27

Db 140 NLWAAQRYGRELRLMSDEFGSFKGLP 166

|||||

RESULT 7

AAW61318

ID AAW61318 standard; Protein; 204 AA.

XX

AC AAW61318;

XX

DT 07-OCT-1998 (first entry)

XX

DE Mutant BCL-XL/BCL-2 associated cell death regulator #3.

XX

KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;

KW serine substituted mutant; apoptosis; cancer; viral infection.

XX

OS Mus sp.

OS Synthetic.

XX

PN WO9817682-A1.

XX

PD 30-APR-1998.

XX

PF 17-OCT-1997; 97WO-US19175.

XX

PR 18-OCT-1996; 96US-0733505.

XX

XX (UNIW) UNIV WASHINGTON.

XX

XX Korsmeyer SJ;

XX

XX WPI; 1998-261422/23.

DR N-PSDB; AAV27836.

DR

PT New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 XX
 XX
 PS Claim 7; Page 60-61; 95pp; English.
 XX
 CC The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX
 XX
 SQ Sequence 204 AA;

Query Match 100.0%; Score 145; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 2.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDFEGSFKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRLMSDFEGSFKGLP 166

RESULT 8
 AAW58832
 ID AAW58832 standard; protein; 204 AA.
 XX
 AC AAW58832;
 XX
 DT 23-JUL-1998 (first entry)
 XX
 DE Murine BAD protein.
 XX
 KW BAD protein; Bcl-XL/Bcl-2 associated cell death regulator; 14-3-3;
 KW serine phosphorylation; post-translational modification; apoptosis;
 KW signal transduction regulator; phosphoserine phosphatase; senescence;
 KW immunodeficiency disease; neurodegenerative disease; infertility;
 KW cancer, viral infection; lymphoproliferative condition; arthritis;
 KW inflammation; autoimmune diseases.

XX Mus sp.
 OS
 PN WO9809643-A1.
 XX
 XX 12-MAR-1998.
 PD
 PF 09-SEP-1997; 97WO-US15871.
 XX
 XX 09-SEP-1996; 96US-0707868.
 PR
 XX (UNIW) UNIV WASHINGTON.
 PA
 XX Korsmeyer SJ;
 PI
 XX WPI; 1998-207049/18.

XX Serine-phosphorylated Bcl-X-L/Bcl-2 Associated cell Death regulator
 PT polypeptide - useful for modulation of apoptosis associated with,
 PT e.g. cancer and immunodeficiency diseases
 XX
 XX Claim 3; Fig 8; 61pp; English.
 PS
 XX This sequence represents a novel serine-phosphorylated protein, BAD
 CC (Bcl-XL/Bcl-2 associated cell death regulator). The serine residue is
 CC phosphorylated in a post-translational modification and allows binding
 CC to the 14-3-3 protein which is a signal transduction regulator.
 CC Modulators of phosphorylated BAD, which act through inhibition/activation
 CC of a phosphoserine phosphatase, are useful for preventing/treating
 CC increased/decreased apoptosis in a cell. The increased apoptosis may
 CC result from immunodeficiency diseases, senescence, neurodegenerative
 CC disease, ischaemic cell death, reperfusion cell death, infertility and
 CC wound-healing. Decreased apoptosis may result from cancer, viral
 CC infection, lymphoproliferative conditions, arthritis, infertility,
 CC inflammation and autoimmune diseases. Measuring the amount of
 CC phosphorylated compared to unphosphorylated BAD polypeptide and/or total
 CC BAD in a cell is useful for determining the apoptotic state of a cell.
 XX
 SQ Sequence 204 AA;

Query Match 100.0%; Score 145; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 2.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDFEGSFKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRLMSDFEGSFKGLP 166

RESULT 9
 AAB70369
 ID AAB70369 standard; protein; 204 AA.

XX
 AC AAB70369;
 XX
 DT 02-MAY-2001 (first entry)
 XX
 XX Longer murine BAD mutant amino acid sequence SEQ ID NO:2.
 DE
 XX Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 KW immunostimulant; neuroprotective; nootropic; antischismic; vulnerary;
 KW cytosolic; antiviral; antiarthritic; antinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.

XX Mus musculus.
 OS Synthetic.
 XX
 PN WO200110888-A1.
 XX
 XX 15-FEB-2001.
 PD
 XX 30-MAY-2000; 2000WO-US11864.
 PF
 XX 28-MAY-1999; 99US-0136783.
 PR
 XX (APOP-) APOPTOSIS TECHNOLOGY INC.
 PA
 XX Zhou X;
 PI
 XX WPI; 2001-138734/14.

XX New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -
 XX

PS Claim 4; Page 148; 157pp; English.

XX The present invention describes an isolated or synthetic polypeptide
CC (I) comprising a less than full length amino acid sequence of a mutant
CC Pcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
CC fragment, which contains amino acid substitutions at Ser118 of a human
CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
CC neurotropic, antiischaemic, vulnerary, cytostatic, antiviral,
CC antiarthritic, antiinflammatory and immunosuppressive activities, and
CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
CC polynucleotides can be used for screening candidate compounds and drugs
CC for activity that promote cell survival or apoptosis. Other uses include
CC inducing or inhibiting apoptosis in a cell. Candidate compounds
CC identified and (mutant) BAD polypeptides are useful in treating
CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
CC death, reperfusion cell death, wound healing, cancer, viral infections,
CC lymphoproliferative conditions, arthritis, infertility, inflammation and
CC autoimmune diseases. The present sequence represents a specifically
CC claimed longer murine BAD mutant amino acid sequence from the present
CC invention.

XX Sequence 204 AA;

Query Match 100.0%; Score 145; DB 22; Length 204;
Best Local Similarity 100.0%; Pred. No. 2.4e-14;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGLP 27
Db 140 NLWAAQRYGRELRRMSDEFGSKGLP 166

RESULT 10
ABR39082
ID ABR39082 standard; Protein; 204 AA.

XX ABR39082;
DT 10-MAY-2003 (first entry)
DE Murine BAD protein SEQ ID NO:4.

KW Murine; BAD; herpes simplex virus; HSV; US3; herpes virus; apoptosis;
KW virucide; infection.

XX Mus musculus.

XX WO2003012049-A2.

XX 13-FEB-2003.

XX 31-JUL-2002; 2002WO-US24177.

XX 31-JUL-2001; 2001US-308929P.

XX (UYCH-) UNIV CHICAGO.

XX Munger J, Roizman B;

XX WPI; 2003-248168/24.

XX N-PSDB; ABZ81201.

XX Inducing apoptosis in a cell infected with herpes simplex virus, HSV,
PT by administering to the cell, a composition comprising an agent that
PT inhibits phosphorylation of pro-apoptotic polypeptide BAD by HSV US3

PS Claim 15; Page 168; 192pp; English.

XX The present invention describes a method (M1) for inducing apoptosis in
CC a cell infected with herpes simplex virus (HSV), which comprises
CC administering to the cell, a composition having an agent that inhibits
CC phosphorylation of pro-apoptotic polypeptide BAD by HSV US3. Also

CC described is a method (M2) for treating a patient infected with HSV, by
CC administering to the patient, a composition comprising a peptide
CC comprising a sequence of 4-100 continuous amino acids of a 168 residue
CC amino acid sequence (see ABR39081), where the peptide comprises Ser112,
CC Ser135, or Ser155, or their combinations. BAD has virucide activity.
CC M1 is useful for inducing apoptosis in a cell infected with HSV, where
CC the cell is in a human. M2 is useful for treating a patient infected
CC with HSV. The present sequence represents murine BAD, which is used in
CC the exemplification of the present invention.

XX Sequence 204 AA;

Query Match 100.0%; Score 145; DB 24; Length 204;
Best Local Similarity 100.0%; Pred. No. 2.4e-14;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGLP 27
Db 140 NLWAAQRYGRELRRMSDEFGSKGLP 166

RESULT 11
AAU00220
ID AAU00220 standard; Protein; 567 AA.

XX AAU00220;

DT 31-MAY-2001 (first entry)

DE Bad-DTRR apoptosis-modifying fusion protein.

XX Mouse; Bad-DTRR: apoptosis; cancer; spinal muscular atrophy;
KW diphtheria toxin receptor binding domain; DTR; neoplasm; tumour;
KW hyper-proliferation; Alzheimer's disease; neurodegenerative disorder;
KW transient ischaemic neuronal injury; stroke; spinal cord injury;
KW Huntington's disease.

XX Chimeric - Mus sp.

OS Chimeric - Corynebacterium diphtheriae.

OS Chimeric - Synthetic.

XX Key Location/Qualifiers

FT Region 3..12 /note= "10x histidine tag"

XX WO200112661-A2.

XX 22-FEB-2001.

XX 15-AUG-2000; 2000WO-US22293.

XX 16-AUG-1999; 99US-0149220.

XX (HARD) HARVARD COLLEGE.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Youle RJ, Liu X, Collier RJ;

XX WPI; 2001-218343/22.

XX N-PSDB; AAS00248.

XX Novel fusion protein for modifying apoptosis in target cell and
PT reducing apoptosis after transient ischaemic neuronal injury, has two
PT domains which targets protein to a cell and modifies apoptotic response
PT of cell

PS Claim 4; Page 59-61; 65pp; English.

XX The sequence represents the amino acid sequence of Bad-DTRR apoptosis-
CC modifying fusion protein comprising Bad gene sequence fused via a short
CC linker to diphtheria toxin translocation domain (DTRR). The
CC functional apoptosis-modifying fusion protein is capable of binding a
CC target cell and integrating into or crossing a cellular membrane of the

CC target cell. The apoptosis-modifying fusion protein comprises at least
 CC two domains: the DTR domain, which targets the fusion protein to the
 CC target cell and the Bcl-XL domain, which modifies an apoptotic response
 CC of the target cell. The fusion protein is useful for modifying
 CC (inhibiting or enhancing) apoptosis in a target cell, such as neuron,
 CC lymphocyte, cancer, neoplasm, macrophage, epithelial, stem, tumour or
 CC hyper-proliferative cell or an adipocyte. It is also useful for reducing
 CC apoptosis in a subject after transient ischemic neuronal injury,
 CC especially spinal cord injury. The fusion protein may be used to treat
 CC various diseases and injury conditions through inhibition or enhancement
 CC of apoptotic cellular response, including neurodegenerative disorders
 CC such as Alzheimer's disease, Huntington's disease, spinal muscular
 CC atrophy, stroke episodes and unregulated cell growth as in tumours and
 CC various cancers. The apoptosis-modifying fusion protein can be delivered
 CC effectively throughout the body and targeted to selective tissue and
 CC cells.

XX SQ Sequence 567 AA;
 Query Match 100.0%; Score 145; DB 22; Length 567;
 Best Local Similarity 100.0%; Pred. No. 7.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 NLWAAQRYGRELRLMSDFEGSFKGLP 27
 DB 161 NLWAAQRYGRELRLMSDFEGSFKGLP 187
 |||||

RESULT 12
 AAB37001
 ID AAB37001 standard; peptide: 26 AA.
 AC AAB37001;
 XX 28-FEB-2001 (first entry)
 DT
 DE Bcl2 polypeptide BH3 domain peptide #1.
 KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.
 XX WO200059526-A1.
 XX 12-OCT-2000.
 XX 06-APR-2000; 2000WO-US09352.
 XX 07-APR-1999; 99US-0128202.
 XX (UYJE-) UNIV JEFFERSON THOMAS.
 XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX WPI; 2000-679325/66.
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 XX Claim 18; Page 17; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH

CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent analogues
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric, or
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX SQ Sequence 26 AA;
 Query Match 95.2%; Score 138; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 3e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDFEGSFKGL 26
 DB 1 NLWAAQRYGRELRLMSDFEGSFKGL 26
 |||||

RESULT 13
 AAB37002
 ID AAB37002 standard; peptide: 26 AA.
 AC AAB37002;
 XX 28-FEB-2001 (first entry)
 DT
 DE Bcl2 polypeptide BH3 domain peptide #2.
 KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.
 XX WO200059526-A1.
 XX 12-OCT-2000.
 XX 06-APR-2000; 2000WO-US09352.
 XX 07-APR-1999; 99US-0128202.
 XX (UYJE-) UNIV JEFFERSON THOMAS.
 XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX WPI; 2000-679325/66.
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 XX Claim 18; Page 17; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where

CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX Sequence 26 AA;
 SQ Query Match 95.2%; Score 138; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 3e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
 DB 1 NLWAAQRYGRELRRMSDFEGSFKGL 26

RESULT 14
 AAB37056
 ID AAB37056 standard; peptide; 27 AA.

XX AAB37056;
 XX
 XX 28-FEB-2001 (first entry)
 XX Bcl2 polypeptide BH3 domain peptide #56.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.
 OS
 XX WO2000059526-A1.
 PN
 XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UJJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer

XX Claim 18; Page 19; 74pp; English.

CC The invention relates to a peptide conjugate having the formula:
 CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX Sequence 27 AA;

Query Match 95.2%; Score 138; DB 21; Length 27;
 Best Local Similarity 100.0%; Pred. No. 3.1e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSFKGL 26
 DB 2 NLWAAQRYGRELRRMSDFEGSFKGL 27

RESULT 15
 AAB37055
 ID AAB37055 standard; peptide; 28 AA.

XX AAB37055;

XX 28-FEB-2001 (first entry)

XX Bcl2 polypeptide BH3 domain peptide #55.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.

OS WO2000059526-A1.

XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UJJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer

XX Claim 18; Page 19; 74pp; English.

PS

XX The invention relates to a peptide conjugate having the formula:
CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the peptide portion of the conjugate. The peptides represent analogues
CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC the Bcl-2 domain of the cell death agonist Bad. The peptide conjugate is
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX

SQ Sequence 28 AA;

Query Match 95.2%; Score 138; DB 21; Length 28;
Best Local Similarity 100.0%; Pred. No. 3.2e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAQRYGRELRRMSDFEGSKGL 26
Db 2 NLWAQRYGRELRRMSDFEGSKGL 27

Search completed: September 15, 2003, 17:22:13
Job time : 38.1857 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:22:21 ; Search time 14.0786 Seconds
(without alignments)
81.144 Million cell updates/sec

Title: US-09-544-664-3

Perfect score: 145

Sequence: 1 NLWAAQRYGRELNRNDEFGSKGLP 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued_patents_AA.*

1: /cgn2.6/ptodata/1/iaa/5A-COMB.pep.*

2: /cgn2.6/ptodata/1/iaa/5B-COMB.pep.*

3: /cgn2.6/ptodata/1/iaa/6A-COMB.pep.*

4: /cgn2.6/ptodata/1/iaa/6B-COMB.pep.*

5: /cgn2.6/ptodata/1/iaa/PTUS-COMB.pep.*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Query %	Length	DB ID	Description
1	145	100.0	204	1	US-08-333-565-2	Sequence 2, Appli
2	145	100.0	204	2	US-08-661-479-2	Sequence 2, Appli
3	145	100.0	204	2	US-08-733-505A-1	Sequence 1, Appli
4	145	100.0	204	2	US-08-733-505A-12	Sequence 13, Appl
5	145	100.0	204	2	US-08-733-505A-13	Sequence 14, Appl
6	145	100.0	204	2	US-08-733-505A-14	Sequence 3, Appli
7	142	97.9	204	2	US-08-717-123-3	Sequence 3, Appli
8	142	97.9	204	4	US-09-375-257-3	Sequence 2, Appli
9	120.5	83.1	166	1	US-08-665-617-2	Sequence 2, Appli
10	120.5	83.1	168	2	US-08-717-123-2	Sequence 1, Appli
11	120.5	83.1	168	3	US-08-985-335-1	Sequence 7, Appli
12	120.5	83.1	168	3	US-08-985-335-7	Sequence 1, Appli
13	120.5	83.1	168	3	US-09-410-372-1	Sequence 7, Appli
14	120.5	83.1	168	3	US-09-410-372-7	Sequence 2, Appli
15	120.5	83.1	168	4	US-09-375-257-2	Sequence 2, Appli
16	113	77.9	23	1	US-08-333-565-10	Sequence 10, Appl
17	113	77.9	23	2	US-08-661-479-10	Sequence 10, Appl
18	102	70.3	59	2	US-08-733-505A-55	Sequence 55, Appl
19	102	70.3	59	2	US-08-733-505A-56	Sequence 56, Appl
20	102	70.3	59	2	US-08-733-505A-57	Sequence 57, Appl
21	102	70.3	59	2	US-08-733-505A-58	Sequence 58, Appl
22	86	59.3	16	1	US-08-333-565-26	Sequence 26, Appl
23	86	59.3	16	2	US-08-661-479-26	Sequence 26, Appl
24	61	42.1	11	2	US-08-733-505A-34	Sequence 34, Appl
25	61	42.1	11	2	US-08-706-741B-69	Sequence 69, Appl
26	61	42.1	11	2	US-08-924-695A-69	Sequence 69, Appl
27	53	36.6	946	3	US-09-074-579-3	Sequence 3, Appli

28 53 36.6 946 3 US-09-388-774-3 Sequence 3, Appli
29 53 36.6 946 4 US-09-546-153-1 Sequence 1, Appli
30 51 35.2 66 2 US-08-867-087B-40 Sequence 40, Appl
31 50.5 34.8 467 4 US-09-252-991A-18296 Sequence 18296, A
32 49 33.8 263 4 US-09-651-856-27 Sequence 27, Appl
33 49 33.8 263 4 US-09-650-855-27 Sequence 27, Appl
34 48.5 33.4 904 4 US-09-328-352-4656 Sequence 4656, Ap
35 46.5 32.1 415 4 US-09-252-991A-31348 Sequence 31348, A
36 46 31.7 322 4 US-09-252-991A-30979 Sequence 30979, A
37 46 31.7 610 4 US-09-252-991A-19594 Sequence 19594, A
38 45.5 31.4 351 4 US-09-252-991A-32466 Sequence 32466, A
39 45.5 31.4 906 4 US-09-252-991A-31458 Sequence 31458, A
40 45 31.0 229 4 US-09-252-991A-23807 Sequence 23807, A
41 45 31.0 303 4 US-09-328-352-5164 Sequence 5164, Ap
42 45 31.0 356 4 US-09-235-103-2 Sequence 2, Appli
43 45 31.0 356 4 US-09-235-103-4 Sequence 4, Appli
44 45 31.0 513 1 US-08-464-340A-2 Sequence 2, Appli
45 45 31.0 513 5 PCT-US94-08449A-2 Sequence 2, Appli

ALIGNMENTS

RESULT 1
US-08-333-565-2
; Sequence 2, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSER: Townsend and Townsend Kourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "Deduced amino acid sequence
; OTHER INFORMATION: of mouse BAD."

Query Match 100.0%; Score 145; DB 1; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.4e-14;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGLP 27
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 Db 140 NLWAAQRYGRELRRMSDEFGSKGLP 166

RESULT 2

US-08-661-479-2
 ; Sequence 2, Application US/08661479
 ; Patent No. 5834209
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, STANLEY J.
 ; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
 ; TITLE OF INVENTION: REGULATOR
 ; NUMBER OF SEQUENCES: 59
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Townsend and Townsend Kourie and Crew
 ; STREET: 379 Lytton Avenue
 ; CITY: Palo Alto
 ; STATE: California
 ; COUNTRY: US
 ; ZIP: 94301
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/661,479
 ; FILING DATE: 11-JUN-1995
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 08/333,565
 ; FILING DATE: 31-OCT-1994
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Smith, William M
 ; REGISTRATION NUMBER: 30,223
 ; REFERENCE/DOCKET NUMBER: 15726A-000700
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (415) 326-2400
 ; TELEFAX: (415) 326-2422
 ; INFORMATION FOR SEQ ID NO: 2:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 204 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; FEATURE:
 ; NAME/KEY: Protein
 ; LOCATION: 1..204
 ; OTHER INFORMATION: /note= "Deduced amino acid sequence
 ; OTHER INFORMATION: Of mouse BAD."
 ;

US-08-661-479-2
 Query Match 100.0%; Score 145; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRRMSDEFGSKGLP 166

RESULT 3

US-08-733-505A-1
 ; Sequence 1, Application US/08733505A
 ; Patent No. 5856445
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, STANLEY J.
 ; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 ; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 ; NUMBER OF SEQUENCES: 60
 ; CORRESPONDENCE ADDRESS:

; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
 ; STREET: 7733 FORSYTH BLVD., SUITE 1400
 ; CITY: ST. LOUIS
 ; STATE: MISSOURI
 ; COUNTRY: USA
 ; ZIP: 63105
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/733,505A
 ; FILING DATE:

; CLASSIFICATION: 530
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: HOLLAND, DONALD R.
 ; REGISTRATION NUMBER: 35,197
 ; REFERENCE/DOCKET NUMBER: 965458
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314) 727-5188
 ; TELEFAX: (314) 727-6092
 ; INFORMATION FOR SEQ ID NO: 1:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 204 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS:
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ;

US-08-733-505A-1

Query Match 100.0%; Score 145; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSKGLP 27
 |||||
 Db 140 NLWAAQRYGRELRRMSDEFGSKGLP 166

RESULT 4

US-08-733-505A-12
 ; Sequence 12, Application US/08733505A
 ; Patent No. 5856445
 ; GENERAL INFORMATION:
 ; APPLICANT: KORSMEYER, STANLEY J.
 ; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 ; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 ; NUMBER OF SEQUENCES: 60
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
 ; STREET: 7733 FORSYTH BLVD., SUITE 1400
 ; CITY: ST. LOUIS
 ; STATE: MISSOURI
 ; ZIP: 63105
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/733,505A
 ; FILING DATE:
 ; CLASSIFICATION: 530
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: HOLLAND, DONALD R.
 ; REGISTRATION NUMBER: 35,197
 ; REFERENCE/DOCKET NUMBER: 965458
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314) 727-5188
 ; TELEFAX: (314) 727-6092
 ; INFORMATION FOR SEQ ID NO: 12:

SEQUENCE CHARACTERISTICS:
 LENGTH: 204 amino acids
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-08-733-505A-12

Query Match 100.0%; Score 145; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGLP 27
 DB 140 NLWAAQRYGRELRLMSDEFGSKGLP 166

RESULT 5

US-08-733-505A-13
 Sequence 13, Application US/08733505A
 Patent No. 5856445

GENERAL INFORMATION:
 APPLICANT: KORSMEYER, STANLEY J.
 TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 NUMBER OF SEQUENCES: 60
 CORRESPONDENCE ADDRESS:

ADDRESSEE: HOWELL & HAFERKAMP, L.C.
 STREET: 7733 FORSYTH BLVD., SUITE 1400
 CITY: ST. LOUIS
 STATE: MISSOURI
 COUNTRY: USA
 ZIP: 63105

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/733,505A

FILING DATE:
 CLASSIFICATION: 530
 ATTORNEY/AGENT INFORMATION:
 NAME: HOLLAND, DONALD R.
 REGISTRATION NUMBER: 35,197

REFERENCE/DOCKET NUMBER: 965458
 TELEPHONE: (314) 727-5188
 TELEFAX: (314) 727-6092

INFORMATION FOR SEQ ID NO: 13:
 LENGTH: 204 amino acids
 TYPE: amino acid
 STRANDEDNESS:

TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-08-733-505A-13

Query Match 100.0%; Score 145; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGLP 27
 DB 140 NLWAAQRYGRELRLMSDEFGSKGLP 166

RESULT 6

US-08-733-505A-14
 Sequence 14, Application US/08733505A
 Patent No. 5856445

GENERAL INFORMATION:
 APPLICANT: KORSMEYER, STANLEY J.

TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
 BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
 NUMBER OF SEQUENCES: 60
 CORRESPONDENCE ADDRESS:

ADDRESSEE: HOWELL & HAFERKAMP, L.C.
 STREET: 7733 FORSYTH BLVD., SUITE 1400
 CITY: ST. LOUIS
 STATE: MISSOURI
 COUNTRY: USA
 ZIP: 63105

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/733,505A

FILING DATE:
 CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:
 NAME: HOLLAND, DONALD R.
 REGISTRATION NUMBER: 35,197
 REFERENCE/DOCKET NUMBER: 965458
 TELECOMMUNICATION INFORMATION:

TELEPHONE: (314) 727-5188
 TELEFAX: (314) 727-6092
 INFORMATION FOR SEQ ID NO: 14:

SEQUENCE CHARACTERISTICS:
 LENGTH: 204 amino acids
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-08-733-505A-14

Query Match 100.0%; Score 145; DB 2; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRLMSDEFGSKGLP 27
 DB 140 NLWAAQRYGRELRLMSDEFGSKGLP 166

RESULT 7

US-08-717-123-3
 Sequence 3, Application US/08717123
 Patent No. 5965703

GENERAL INFORMATION:
 APPLICANT: Horne, William A.

ADDRESSEE: Oltersdorf, Tilman
 TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
 Acids and Methods of Use
 NUMBER OF SEQUENCES: 15

CORRESPONDENCE ADDRESS:
 ADDRESSEE: Campbell and Flores
 STREET: 4370 La Jolla Village Drive, Suite 700
 CITY: San Diego
 STATE: California
 COUNTRY: United States
 ZIP: 92122

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/717,123
 FILING DATE: 20-SEP-1996
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Campbell, Cathryn A.
 REGISTRATION NUMBER: 31,815

REFERENCE/DOCKET NUMBER: P-ID 1929
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
TOPOLOGY: linear
US-08-717-123-3

Query Match 97.9%; Score 142; DB 2; Length 204;
Best Local Similarity 96.3%; Pred. No. 4.1e-14;
Matches 26; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSF-KGLP 27
Db 140 NLWAAQRYGRELRRMTDFEGSF-KGLP 166

RESULT 8
US-09-375-257-3
Sequence 3, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:
APPLICANT: Horne, William A.
TITLE OF INVENTION: HUMAN RAD POLYPEPTIDES, ENCODING NUCLEIC
FILE REFERENCE: 480140.428D1
CURRENT APPLICATION NUMBER: US/09/375,257
CURRENT FILING DATE: 1999-08-16
NUMBER OF SEQ ID NOS: 15
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 3
LENGTH: 204
TYPE: PRT
ORGANISM: Mus musculus
US-09-375-257-3

Query Match 97.9%; Score 142; DB 4; Length 204;
Best Local Similarity 96.3%; Pred. No. 4.1e-14;
Matches 26; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDFEGSF-KGLP 27
Db 140 NLWAAQRYGRELRRMTDFEGSF-KGLP 166

RESULT 9
US-08-665-617-2
Sequence 2, Application US/08665617
Patent No. 5663316
GENERAL INFORMATION:
APPLICANT: Xudong, Yin
TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Saliwanchik & Saliwanchik
STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: Florida
COUNTRY: USA
ZIP: 32606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/665,617
FILING DATE:
CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:
NAME: Saliwanchik, David R.
REGISTRATION NUMBER: 31,794
REFERENCE/DOCKET NUMBER: CL-8
TELEPHONE: (352) 375-8100
TELEFAX: (352) 372-5800
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 166 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-665-617-2

Query Match 83.1%; Score 120.5; DB 1; Length 166;
Best Local Similarity 89.3%; Pred. No. 6.2e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDFEGSF-KGLP 27
Db 101 NLWAAQRYGRELRRMSDFVDSFKKGLP 138

RESULT 10
US-08-717-123-2
Sequence 2, Application US/08717123
Patent No. 5965703
GENERAL INFORMATION:
APPLICANT: Horne, William A.
TITLE OF INVENTION: Human RAD Polypeptides, Encoding Nucleic
TITLE OF INVENTION: Acids and Methods of Use
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell and Flores
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: United States
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/717,123
FILING DATE: 20-SEP-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-ID 1929
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-717-123-2

Query Match 83.1%; Score 120.5; DB 2; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDFEGSF-KGLP 27
Db 103 NLWAAQRYGRELRRMSDFVDSFKKGLP 130

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RESULT 11
US-08-985-335-1
; Sequence 1, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
; US-08-985-335-1

Query Match 83.1%; Score 120.5; DB 3; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRLRMSDEFGSF-KGLP 27
Db 103 NLWAAQRYGRLRMSDEFGSF-KGLP 130

RESULT 12
US-08-985-335-7
; Sequence 7, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9

```

```

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; US-08-985-335-7

Query Match 83.1%; Score 120.5; DB 3; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRLRMSDEFGSF-KGLP 27
Db 103 NLWAAQRYGRLRMSDEFGSF-KGLP 130

RESULT 13
US-09-410-372-1
; Sequence 1, Application US/09410372
; Patent No. 6281334
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:

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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
;
US-09-410-372-1

Query Match      83.1%; Score 120.5; DB 3; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDEFEQSF-KGLP 27
Db 103 NLWAAQRYGRELRRMSDEFEVDSFKKGLP 130

RESULT 15
US-09-375-257-2
; Sequence 2, Application US/09375257
; Patent No. 6504022
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.42851
; CURRENT APPLICATION NUMBER: US/09/375,257
; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: Fast-SEQ for Windows Version 4.0
; SEQ ID NO: 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
;
US-09-375-257-2

Query Match      83.1%; Score 120.5; DB 4; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDEFEQSF-KGLP 27
Db 103 NLWAAQRYGRELRRMSDEFEVDSFKKGLP 130

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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
;
US-09-410-372-1

Query Match      83.1%; Score 120.5; DB 3; Length 168;
Best Local Similarity 89.3%; Pred. No. 6.3e-11;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDEFEQSF-KGLP 27
Db 103 NLWAAQRYGRELRRMSDEFEVDSFKKGLP 130

RESULT 14
US-09-410-372-7
; Sequence 7, Application US/09410372
; Patent No. 6281334
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSER: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
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GenCore version 5.1.6
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OM protein - protein search, using sw model

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Title: US-09-544-664-3

Perfect score: 145
Sequence: 1 NLWAAQRYGRELRLMSDFEGSKGLP 27

Scoring table: BLOSUM62
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Searched: 541936 seqs, 145912426 residues

Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:*

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12: /cgn2_6/ptodata/1/pubaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	142	97.9	204	9	US-09-922-378-3
2	142	97.9	204	14	US-10-066-179-3
3	120.5	83.1	168	9	US-09-922-378-2
4	120.5	83.1	168	9	US-09-994-657-1
5	120.5	83.1	168	9	US-09-994-657-7
6	120.5	83.1	168	14	US-10-066-179-2
7	114	78.6	25	15	US-10-059-261-258
8	71	49.0	15	15	US-10-174-105A-147
9	53	36.6	946	9	US-09-828-423-3
10	50	34.5	215	15	US-10-156-761-9145
11	47	32.4	35	15	US-10-092-750-1
12	47	32.4	138	15	US-10-092-750-241
13	46	31.7	682	12	US-10-238-075-1077
14	45.5	31.4	852	9	US-09-752-639-153
15	45.5	31.4	852	10	US-09-984-198-153

16	45	31.0	513	12	US-10-199-869-4	Sequence 4, Appli
17	45	31.0	513	14	US-10-143-002-2	Sequence 2, Appli
18	45	31.0	513	15	US-10-325-891-2	Sequence 2, Appli
19	44.5	30.7	144	12	US-09-903-190-160	Sequence 160, App
20	44.5	30.7	146	15	US-10-121-757B-12	Sequence 12, Appl
21	44.5	30.7	160	10	US-09-738-973-435	Sequence 435, App
22	44.5	30.7	160	10	US-09-854-133-435	Sequence 435, App
23	44.5	30.7	160	15	US-10-144-649A-435	Sequence 435, App
24	44.5	30.7	309	15	US-10-102-806-649	Sequence 649, App
25	44.5	30.7	334	10	US-09-794-715A-8	Sequence 8, Appli
26	44.5	30.7	334	12	US-10-286-581-8	Sequence 8, Appli
27	44.5	30.7	334	15	US-10-046-924-8	Sequence 8, Appli
28	44	30.3	81	9	US-09-925-297-560	Sequence 560, App
29	44	30.3	272	15	US-10-156-761-11541	Sequence 11541, A
30	44	30.3	426	9	US-09-815-242-5704	Sequence 5704, Ap
31	44	30.3	699	14	US-10-008-355-8	Sequence 8, Appli
32	44	30.3	705	9	US-09-815-242-12463	Sequence 12463, A
33	44	30.3	712	14	US-10-008-355-2	Sequence 2, Appli
34	44	30.3	877	12	US-10-369-294-20	Sequence 20, Appl
35	43.5	30.0	543	15	US-10-156-761-13485	Sequence 13485, A
36	43	29.7	213	9	US-09-843-846-18	Sequence 18, Appl
37	43	29.7	232	10	US-09-881-752A-238	Sequence 238, App
38	43	29.7	270	11	US-09-934-455-162	Sequence 162, App
39	43	29.7	380	9	US-09-149-045-2	Sequence 2, Appli
40	43	29.7	380	15	US-10-166-359-2	Sequence 2, Appli
41	43	29.7	380	15	US-10-166-113-2	Sequence 2, Appli
42	43	29.7	380	15	US-10-166-357-2	Sequence 2, Appli
43	43	29.7	380	15	US-10-166-372-2	Sequence 2, Appli
44	43	29.7	380	15	US-10-184-722-3	Sequence 3, Appli
45	43	29.7	380	15	US-10-251-385-62	Sequence 62, Appl

ALIGNMENTS

RESULT 1
US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PPT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 97.9%; Score 142; DB 9; Length 204;
Best Local Similarity 96.3%; Pred. No. 2.5e-13;
Matches 26; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRELRLMSDFEGSKGLP 27
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Db 140 NLWAAQRYGRELRLMTDFEGSKGLP 166

RESULT 2
US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE

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; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066.179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-066-179-3

Query Match          97.9%; Score 142; DB 14; Length 204;
Best Local Similarity 96.3%; Pred. No. 2.5e-13;
Matches 26; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 NLWAAQRYGRELRLMSDEFGSKGLP 27
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Db       140 NLWAAQRYGRELRLMTDEFGSKGLP 166

RESULT 3
US-09-922-378-2
; Sequence 2, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-922-378-2

Query Match          83.1%; Score 120.5; DB 9; Length 168;
Best Local Similarity 89.3%; Pred. No. 3.2e-10;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY      1 NLWAAQRYGRELRLMSDEFGSP-KGLP 27
        |||||
Db       103 NLWAAQRYGRELRLMSDEFVDSFKKGLP 130

RESULT 4
US-09-894-657-1
; Sequence 1, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
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; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166

INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: SYNORAE01
CLONE: 358673
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-894-657-1

Query Match          83.1%; Score 120.5; DB 9; Length 168;
Best Local Similarity 89.3%; Pred. No. 3.2e-10;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY      1 NLWAAQRYGRELRLMSDEFGSP-KGLP 27
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Db       103 NLWAAQRYGRELRLMSDEFVDSFKKGLP 130

RESULT 5
US-09-894-657-7
; Sequence 7, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
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; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 7:
US-09-894-657-7

Query Match 83.1%; Score 120.5; DB 9; Length 168;
Best Local Similarity 89.3%; Pred. No. 3.2e-10;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDEFGSF-KGLP 27
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Db 103 NLWAAQRYGRELRRMSDEFGSF-KGLP 130

RESULT 6

US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-066-179-2

Query Match 83.1%; Score 120.5; DB 14; Length 168;
Best Local Similarity 89.3%; Pred. No. 3.2e-10;
Matches 25; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

QY 1 NLWAAQRYGRELRRMSDEFGSF-KGLP 27
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Db 103 NLWAAQRYGRELRRMSDEFGSF-KGLP 130

RESULT 7

US-10-059-261-258
; Sequence 258, Application US/10059261
; Publication No. US20030077826A1
; GENERAL INFORMATION:
; APPLICANT: EDELMAN, LENA
; APPLICANT: JACOPOT, ETIENNE DANIEL FRANCOIS
; APPLICANT: BRIAND, JEAN-PAUL
; TITLE OF INVENTION: CHIMERIC MOLECULES CONTAINING A MODULE ABLE TO TARGET
; TITLE OF INVENTION: SPECIFIC CELLS AND A MODULE REGULATING THE APOPTOGENIC
; TITLE OF INVENTION: FUNCTION OF THE PERMEABILITY TRANSITION PORE COMPLEX
; TITLE OF INVENTION: (PTPC)
; FILE REFERENCE: 03495.0216
; CURRENT APPLICATION NUMBER: US/10/059,261
; CURRENT FILING DATE: 2002-08-29
; PRIOR APPLICATION NUMBER: 60/265,594
; PRIOR FILING DATE: 2001-02-02
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 258
; LENGTH: 25
; TYPE: PRT
; ORGANISM: Unknown Organism

; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: TOX peptide
US-10-059-261-258

Query Match 78.6%; Score 114; DB 15; Length 25;
Best Local Similarity 91.7%; Pred. No. 4.1e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRELRRMSDEFGSF 24
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Db 1 NLWAAQRYGRELRRMSDEFGSF 24

RESULT 8

US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US20030068652A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECI
; FILE REFERENCE: CST-138 CIP3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147

Query Match 49.0%; Score 71; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.00058;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 GRELRMSDEFGS 22
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Db 1 GRELRMSDEFGS 14

RESULT 9

US-09-828-423-3
; Sequence 3, Application US/09828423
; Patent No. US20020099178A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Guegler, Karl J.
; APPLICANT: Patterson, Chandra
; TITLE OF INVENTION: GROWTH-ASSOCIATED TRYPsin-TYPE
; INHIBITOR HEAVY CHAIN PRECURSOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:18:16 ; Search time 12.15 seconds
(without alignments)
213.708 Million cell updates/sec

Title: US-09-544-664-3
Perfect score: 145
Sequence: 1 NLWAAQRYGRLRRMSDEFGSFKGLP 27

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_76.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	DB ID	Description
1	145	100.0	204	2	A55671	bad protein - mouse
2	61	42.1	946	2	JC5575	inter-alpha-trypsi
3	60	41.4	946	2	S54354	inter-alpha-inhibi
4	53	36.6	223	2	D70760	hypothetical prote
5	53	36.6	946	1	IVHU2	inter-alpha-trypsi
6	52	35.9	370	2	S38185	2-dehydro-3-deoxy-
7	51	35.2	232	2	A42095	floral homeotic pr
8	50	34.5	374	2	C84338	spermidine/putresc
9	50	34.5	516	2	A96753	probable threonine
10	49	33.8	263	2	A64807	endonuclease VIII
11	49	33.8	263	2	A85572	hypothetical prote
12	49	33.8	263	2	C90721	hypothetical prote
13	49	33.8	263	2	AD0590	endonuclease VIII,
14	49	33.8	453	2	E83317	conserved hypothet
15	48.5	33.4	134	2	S40376	ig kappa chain - h
16	48.5	33.4	314	2	T02975	annexin P35 - maiz
17	48	33.1	206	2	C36365	transforming prote
18	48	33.1	220	2	F72289	oxidoreductase, so
19	48	33.1	526	2	T08545	threonine synthase
20	47.5	32.8	779	2	B81287	hypothetical prote
21	47	32.4	597	2	G82308	oxaloacetate decar
22	47	32.4	967	2	F82668	oxoglutarate dehyd
23	47	32.4	5138	2	B96695	hypothetical prote
24	46.5	32.1	314	2	T02961	annexin P33 - maiz
25	46.5	32.1	435	2	A44308	Antho-RFamide prec
26	46.5	32.1	1140	2	T09486	hypothetical prote
27	46	31.7	399	2	T35440	probable polyamine
28	46	31.7	497	2	C98318	alkaline proteinase
29	46	31.7	1164	2	T24806	hypothetical prote

ALIGNMENTS

RESULT 1

A55671
bad protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999
C:Accession: A55671
R:Yang, E.; Zha, J.; Jockel, J.; Boise, L.H.; Thompson, C.B.; Korsmeyer, S.J.
Cell 80, 285-291, 1995
A:Title: Bad, a heterodimeric partner for Bcl-x-L and Bcl-2, displaces Bax and promot
A:Reference number: A55671; MUID:95136361; PMID:7834748
A:Accession: A55671
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-204 <YAN>
A:Cross-references: GB:L37296; NID:g639778; PIDN:AAA64465.1; PID:g639779
C:Keywords: heterodimer

Query Match 100.0%; Score 145; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1e-13;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRLRRMSDEFGSFKGLP 27
Db 140 NLWAAQRYGRLRRMSDEFGSFKGLP 166
|||||

RESULT 2

JC5575
inter-alpha-trypsin inhibitor heavy chain 2 - golden hamster
C:Species: Mesocricetus auratus (golden hamster)
C:Date: 23-Sep-1997 #sequence_revision 23-Sep-1997 #text_change 20-Jun-2000
C:Accession: JC5575; PC4485
R:Nakatani, T.; Suzuki, Y.; Yamamoto, T.; Sinothara, H.
J. Biochem. 122, 71-82, 1997
A:Title: Molecular cloning and sequencing of cDNAs encoding three heavy-chain precurs
sin inhibitor heavy chain family
A:Reference number: JC5574; MUID:97420688; PMID:9276673
A:Accession: JC5575
A:Molecule type: mRNA
A:Residues: 1-946 <NAK>
A:Cross-references: DBJ:D89286; NID:g1694689; PIDN:BAAI3939.1; PID:g1694690
A:Experimental source: liver
A:Accession: PC4485
A:Molecule type: protein
A:Residues: 55-64;140-146;151-156;424-447;500-528;577-605 <NA2>
C:Comment: In the plasma three inter-alpha-trypsin inhibitor heavy chains 1, 2 and 3
that the complexes play important role for pancreatic cancer.
C:Superfamily: inter-alpha-trypsin inhibitor complex component II
F:261-264,717-916/Disulfide bonds: #status predicted

Query Match 42.1%; Score 61; DB 2; Length 946;
Best Local Similarity 37.0%; Pred. No. 1;

A:Title: Complementary DNA and derived amino acid sequence of the precursor of one of
A:Reference number: S00346; MUID:88152237; PMID:2450046
A:Accession: S00346
A:Molecule type: mRNA
A:Residues: 1-946 <GB>
A:Cross-references: EMBL:X07173
A:Experimental source: liver
A:Note: part of this sequence, including the amino end of the mature protein, was con-
sisting occurs at the carboxyl as well as the amino end to produce the mature protein
A:Note: due to a double frameshift, the nucleic acid sequence of codons 363-372 is in
A:Note: in one clone, a T is lacking from codon 716; this clone could code for a prot
R:Schreimuelier, T.; Hochstrasser, K.; Reisinger, P.W.M.; Wachter, E.; Gebhard, W.
Biol. Chem. Hoppe-Seyler 368, 963-970, 1987
A:Title: cDNA cloning of human inter-alpha-trypsin inhibitor discloses three differen
A:Reference number: S09064; MUID:38024442; PMID:3663330
A:Accession: S09064
A:Molecule type: mRNA
A:Residues: 265,'RR',268-284,'D',286-946 <SCH>
A:Note: this sequence has been revised in reference S00346
R:Salier, J.P.; Diarra-Mehrpour, M.; Sebboue, R.; Bourguignon, J.; Benarous, R.; Ohku
Proc. Natl. Acad. Sci. U.S.A. 84, 8272-8276, 1987
A:Title: Isolation and characterization of cDNAs encoding the heavy chain of human in
A:Reference number: A39967; MUID:89068576; PMID:2446322
A:Accession: A39967
A:Molecule type: mRNA
A:Residues: 384-673,'A',675-704,'S',706-728,'D',730,'A',732-865 <SAL>
A:Cross-references: GB:M18193; GB:J03013; NID:G338222; PIDN:AA60558.1; PID:G553647
A:Experimental source: liver
R:Salier, J.P.; Diarra-Mehrpour, M.; Sebboue, R.; Bourguignon, J.; Martin, J.P.
Biol. Chem. Hoppe-Seyler 369(suppl.), 15-18, 1988
A:Title: Human inter-alpha-trypsin inhibitor. Isolation and characterization of heavy
A:Reference number: S00632; MUID:89076497; PMID:2462430
A:Accession: S00632
A:Molecule type: mRNA
A:Residues: 384-673,'A',675-704,'S',706-728,'D',730,'A',732-766 <SA2>
A:Cross-references: GB:M3033; NID:G186589; PIDN:AA559195.1; PID:G186590
R:Englind, J.J.; Thøgersen, I.B.; Pizzo, S.V.; Salvesen, G.
J. Biol. Chem. 264, 15975-15981, 1989
A:Title: Analysis of inter-alpha-trypsin inhibitor and a novel trypsin inhibitor, pre
A:Reference number: A92736; MUID:89380192; PMID:2476436
A:Accession: B34245
A:Molecule type: protein
A:Residues: 55-74 <ENG>
R:Maiki, N.; Baidyuk, M.; Maes, P.; Capon, C.; Mizon, C.; Han, K.K.; Tartar, A.; Fou
Biol. Chem. Hoppe-Seyler 373, 1009-1018, 1992
A:Title: The heavy chains of human plasma inter-alpha-trypsin inhibitor: their isolat
A:Reference number: S28928; MUID:93039735; PMID:1384548
A:Accession: S28929
A:Molecule type: protein
A:Residues: 55-64 <MAL>
R:Wisniewski, H.G.; Burgess, W.H.; Oppenheim, J.D.; Vilcek, J.
Biochemistry 33, 7423-7429, 1994
A:Title: TSG-6, an arthritis-associated hyaluronan binding protein, forms a stable co
A:Reference number: A53642; MUID:94271799; PMID:7516184
A:Accession: C53642
A:Molecule type: protein
A:Residues: 55-64 <WIS>
C:Comment: Inter-alpha-trypsin inhibitor is a complex of three proteins, each derivin
C:Note: This protein is a heterodimer of heavy and light chains.
C:Genetics:
A:Gene: GDB:ITTH2
A:Cross-references: GDB:120108; OMIM:146640
A:Map position: 10p15-10p15
C:Superfamily: inter-alpha-trypsin inhibitor complex component II
C:Keywords: carboxylglutamic acid; glycoprotein; heterodimer; serine proteinase inhibi
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-54/Domain: propeptide #status predicted <PRO>
F:55-698/Product: inter-alpha-trypsin inhibitor heavy chain 2 #status predicted <VAR>
F:96-445/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:118,671/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:282,283/Modified site: gamma-carboxylglutamic acid (Glu) #status predicted
F:421,422,423/Binding site: calcium (Asp, Gly, Asp) #status predicted

Db 80 DLEAAQFVALRLKLSDELKG 100

RESULT 7

A42095

floral homeotic protein APETALA3 (AP3) - Arabidopsis thaliana

N;Alternate names: homeotic protein APETALA3; MADS-box regulatory protein AP3

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 04-Mar-1993 #sequence_revision 18-Nov-1994 #text_change 21-Jul-2000

C;Accession: A42095; S52633; T47593

R;Jack, T.; Brockman, L.L.; Meyerowitz, E.M.

Cell 68, 683-697, 1992

A;Title: The homeotic gene APETALA3 of Arabidopsis thaliana encodes a MADS box and is

A;Reference number: A42095; MUID:92154682; PMID:1346756

A;Accession: A42095

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-232 <JAC>

A;Cross-references: GB:M86357; NID:gl66607; PIDN:AAA32740.1; PID:gl66608

A;Experimental source: petals, stamens

A;Note: sequence extracted from NCBI backbone (NCBIIN:82520, NCBIFP:82521)

R;Okamoto, H.; Yano, A.; Shiraishi, H.; Okada, K.; Shimura, Y.

Plant Mol. Biol. 26, 445-472, 1994

A;Title: Genetic complementation of a floral homeotic mutation, ap3, with an Ara

A;Reference number: S52633; MUID:95036018; PMID:7948893

A;Accession: S52633

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-83 <ORA>

A;Cross-references: GB:D21125

R;Glockner, H.; Mewes, H.W.; Lemcke, K.; Mayer, K.F.X.; Quetier, F.; Salanoubat M.M.W.

submitted to the Protein Sequence Database, March 2000

A;Reference number: Z24469

A;Accession: T47593

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-232 <ELO>

A;Cross-references: EMBL:AL132971

A;Experimental source: cultivar Columbia; BAC clone T12E18

C;Genetics:

A;Map position: 3

A;Introns: 63/2; 85/3; 106/2; 139/3; 153/3; 168/3

A;Note: T12E18.30

C;Superfamily: transcription factor sqwa; serum response factor DNA-binding domain h

C;Keywords: DNA binding; nucleus: transcription regulation

F;2-57/Domain: serum response factor DNA-binding domain homology <SRF>

Query Match 35.2%; Score 51; DB 2; Length 232;

Best Local Similarity 44.4%; Pred. No. 7;

Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRMSEFECSFK 24

||| :||| :||| :||| :|||

Db 107 QRLGCELDLDTQLRLLEDEMTFK 133

RESULT 8

C84338

spermidine/putrescine ABC transporter [imported] - Halobacterium sp. NRC-1

C;Species: Halobacterium sp. NRC-1

C;Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001

C;Accession: C84338

R;Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky

; Leithauser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Ja

Jung, K.H.; Alam, M.; Freitas, T.

Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000

A;Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.;

A;Title: Genome sequence of Halobacterium species NRC-1

A;Reference number: A84160; MUID:20504483; PMID:11016950

A;Accession: C84338

A;Status: preliminary

A;Molecule type: DNA

```

Query Match          33.8%; Score 49; DB 2; Length 263;
Best Local Similarity 47.4%; Pred. No. 16;
Matches          9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY      9  GRLRRMSDEFGSFKGLP 27
          | : | | : | | : | | |
Db       4  GPEIRRAADNLEAAIKGRP 22

RESULT 11
A85572
hypothetical protein nei [imported] - Escherichia coli (strain O157:H7, substrain EDL
C: Species: Escherichia coli
C: Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 17-May-2002
C: Accession: A85572
R: Perrona, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; May
iller, L.; Grobeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apoda
Nature 409, 529-533, 2001
A: Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A: Reference number: A85480; PMID:11206551
A: Accession: A85572
A: Status: preliminary
A: Molecule type: DNA
A: Residues: 1-263 <SPC>
A: Cross-references: GB:AE005174; NID:q12513625; PIDN:AG55037.1; GSPDB:GN00145; UWGP
A: Experimental source: strain O157:H7, substrain EDL933
C: Genets:
A: Gene: nei
C: Superfamily: formamidopyrimidine-DNA glycosidase

Query Match          33.8%; Score 49; DB 2; Length 263;
Best Local Similarity 47.4%; Pred. No. 16;
Matches          9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY      9  GRLRRMSDEFGSFKGLP 27
          | : | | : | | : | | |
Db       4  GPEIRRAADNLEAAIKGRP 22

```

```

RESULT 12
C90721
hypothetical protein Ecs0739 [imported] - Escherichia coli (strain O157:H7, substrain
C;Species: Escherichia coli
C;Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 17-May-2002
C;Accession: C90721
R;Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shibata, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A;Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and 9
A;Reference number: A99629; MWID:21156231; PMID:11258796
A;Accession: C90721
A>Status: preliminary
A:Molecule type: DNA
A;Residues: 1-263 <HAY>
A;Cross-references: GB:BA000007; PIDN:BA834162.1; PID:gl3360197; GSPDB:GN00154
A;Experimental source: strain O157:H7, substrain RMD 0509952
C;Genetics:
A;Gene: PCs0739
C;Superfamily: formamidopyrimidine-DNA glycosylase

Query Match          33.8%; Score 49; DB 2; Length 263;
Best Local Similarity 47.4%; Pred. No. 16;
Matches 9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY      9 GRELRMSDEFEGSKGLP 27
         ||||| : | : | : |
Db       4 GPETRRADNLAAIKGKP 22

RESULT 13
AD0590
endonuclease VIII, DNA N-glycosylase with an AP lyase activity STY0771 [imported] - S
```


C:Species: *Salmonella enterica* subsp. *enterica* serovar Typhi
 A:Note: this species has also been called *Salmonella typhi*
 C:Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov-2002
 C:Accession: AD0590
 R:Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher, T.; Connor, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar, S.; Moulie, S.; O'Gaora, P.
 Nature 413, 848-852, 2001
 A:Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; A:Title: Complete genome sequence of a multiple drug resistant *Salmonella enterica* serovar
 A:Reference number: AB0502; MUID:21534947; PMID:11677608
 A:Accession: AD0590
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-263 <PAR>
 A:Cross-references: GB:AL513382; PIDN:CAD05190.1; PID:g16501960; GSPDB:GN00176
 C:Genetics:
 A:Gene: STY0771
 C:Superfamily: formamidopyrimidine-DNA glycosylase

Query Match 33.8%; Score 49; DB 2; Length 263;
 Best Local Similarity 47.4%; Pred. No. 16;
 Matches 9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 9 GRELRMSDEFGSGFKGLP 27
 DB 4 GPEIRADNLEAIAKGP 22
 ||:|||||:|||||

RESULT 14

E83517
 conserved hypothetical protein PA1031 [imported] - *Pseudomonas aeruginosa* (strain PA01)
 C:Species: *Pseudomonas aeruginosa*
 C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
 C:Accession: E83517
 R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.; Berman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim, J.; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000
 A:Title: Complete genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic pathogen
 A:Reference number: AB2950; MUID:20437337; PMID:10984043
 A:Accession: E83517
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-453 <STO>
 A:Cross-references: GB:AE004535; GB:AE004091; NID:g9946936; PIDN:AA04420.1; GSPDB:GN00176
 A:Experimental source: strain PA01
 C:Genetics:
 A:Gene: PA1031

Query Match 33.8%; Score 49; DB 2; Length 453;
 Best Local Similarity 55.6%; Pred. No. 28;
 Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;

QY 3 WAAQRYGR--ELRRMSDE 18
 DB 65 WASERGREELRLRLASE 82
 ||:|||||:|||||

RESULT 15

S40376
 Ig kappa chain - human
 C:Species: *Homo sapiens* (man)
 C:Date: 06-Mar-1994 #sequence_revision 26-May-1995 #text_change 21-Jan-2000
 C:Accession: S40376
 R:Klein, R.; Jaenichen, R.; Zachau, H.G.
 Eur. J. Immunol. 23, 3248-3271, 1993
 A:Title: Expressed human immunoglobulin chi genes and their hypermutation.
 A:Reference number: S40312; MUID:94080891; PMID:8258341
 A:Accession: S40376
 A>Status: preliminary; translation not shown
 A:Molecule type: mRNA
 A:Residues: 1-134 <KLE>

A:Cross-references: EMBL:X72486; NID:g441440; PIDN:CAA51154.1; PID:g441441
 C:Superfamily: immunoglobulin V region; immunoglobulin homology
 C:Keywords: heterotrimer; immunoglobulin
 F:34-I13/Domain: immunoglobulin homology <IMM>

Query Match 33.4%; Score 48.5; DB 2; Length 134;
 Best Local Similarity 38.2%; Pred. No. 9.3;
 Matches 13; Conservative 1; Mismatches 9; Indels 11; Gaps 1;
 QY 3 WAAQRYGRELRM-----SDEFGSGFKG 25
 |||||:|||||
 DB 58 WFRQREGSPRLIYNVSKRDSGVDFSGSG 91

Search completed: September 15, 2003, 17:27:01
 Job time : 13.15 secs

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:55 ; Search time 6.36429 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-3
Perfect score: 145
Sequence: 1 NLWAQRYGRLRMSDFEGSKGLP 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	145	100.0	204	1 BAD_MOUSE	O61337 mus musculus
2	145	100.0	205	1 BAD_RAT	O35147 rattus norv
3	120.5	83.1	188	1 BAD_HUMAN	O92934 homo sapien
4	61	42.1	946	1 ITH2_MESAU	P97279 mesocricetu
5	60	41.4	946	1 ITH2_MOUSE	O61703 mus musculu
6	53	36.6	946	1 ITH2_HUMAN	P19823 homo sapien
7	52	35.9	370	1 AROG_YEAST	P32449 saccharomyc
8	51	35.2	232	1 AP3_ARATH	P35632 arabidopsis
9	50	34.5	851	1 CE05_MOUSE	O8k2h3 mus musculu
10	49.5	34.1	506	1 MATK_LEDPA	O62392 ledum palus
11	49.5	34.1	506	1 MATK_RHOFR	O62984 rhododendro
12	49.5	34.1	506	1 MATK_RHOVS	O62991 rhododendro
13	49	33.8	262	1 ENB8_ECO57	O8x9c6 escherichia
14	49	33.8	262	1 ENB8_ECOLI	P50465 escherichia
15	49	33.8	262	1 ENB8_SALTI	O8z8d2 salmonella
16	49	33.8	262	1 ENB8_SALTY	O8z8u6 salmonella
17	49	33.8	453	1 FMOC_PSEAE	O914t3 pseudomonas
18	48	33.1	205	1 RAS3_RHREA	P22280 rhizomucor
19	48	33.1	220	1 6PGL_THEMA	O9x0n8 thermotoga
20	48	33.1	519	1 THRC_SOLTU	O9mt28 solanum tub
21	48	33.1	526	1 THRC_ARATH	O9s7b5 arabidopsis
22	47	32.4	198	1 BIM_HUMAN	O43521 homo sapien
23	46.5	32.1	429	1 FMR2_ATEL	O16419 anthopleura
24	46.5	32.1	435	1 FMR1_ATEL	P10194 anthopleura
25	46.5	32.1	855	1 XAB2_MOUSE	O9ddcd mus musculu
26	46.5	32.1	855	1 XAB2_RAT	O99pk0 rattus norv
27	46	31.7	738	1 SEC6_DROME	O9v8k2 drosophila
28	46	31.7	1378	1 RPOB_CAWJE	O46124 campylobact
29	45.5	31.4	287	1 PRFA_POLPE	P21359 polyorchis
30	45.5	31.4	334	1 FMRA_CALPA	O01133 calliactis
31	45.5	31.4	507	1 MATK_LOIPR	O47169 loiseleuria
32	45.5	31.4	855	1 XAB2_HUMAN	O9hcs7 homo sapien
33	45	31.0	328	1 SNF4_KLUJA	O9p869 kluyveromyc

RESULT 1

BAD_MOUSE

ID	BAD_MOUSE	STANDARD	PRT	204 AA.
AC	Q61337			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	28-FEB-2003 (Rel. 41, Last annotation update)			
DE	Bcl-2-antagonist of cell death (BAD) (Bcl-2 binding component			
DE	6) (Bcl-xL/Bcl-2 associated death promoter).			
GN	BAD OR BHC6.			
OS	Mus musculus (Mouse).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.			
OX	NCBI_TaxID=10090;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Brain, and Thymus;			
RC	MEDLINE=95136361; PubMed=7834748;			
RA	Yang E., Zha J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;			
RT	"Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and			
RT	promotes cell death."			
RL	Cell 80:285-291(1995).			
RN	[2]			
RP	PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.			
RX	MEDLINE=98022383; PubMed=9381178;			
RA	Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;			
RT	"Interleukin-3-induced phosphorylation of BAD through the protein			
RT	kinase Akt."			
RL	Science 278:687-689(1997).			
RN	[3]			
RP	MUTAGENESIS OF SERINE RESIDUES.			
RX	MEDLINE=20403302; PubMed=10949026;			
RA	Datta S.R., Katsov A., Hu L., Petros A., Fesik S.W., Yaffe M.B.,			
RA	Greenberg M.E.;			
RT	"14-3-3 proteins and survival kinases cooperate to inactivate BAD by			
RT	BH3 domain phosphorylation."			
RL	Mol. Cell 6:41-51(2000).			
CC	-I- FUNCTION: Promotes cell death. Successfully competes for the			
CC	binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level			
CC	of heterodimerization of these proteins with BAX. Can reverse the			
CC	death repressor activity of Bcl-x(L), but not that of Bcl-2.			
CC	Appears to act as a link between growth factor receptor signaling			
CC	and the apoptotic pathways.			
CC	-I- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-			
CC	x(L), Bcl-2 and Bcl-w. Also binds protein Sl00A10 (By similarity).			
CC	The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.			
CC	-I- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon			
CC	phosphorylation, locates to the cytoplasm.			
CC	-I- DOMAIN: Intact BH3 domain is required by BIK, BID, BAX, BAD AND			
CC	BAX for their pro-apoptotic activity and for their interaction			
CC	with anti-apoptotic members of the Bcl-2 family.			
CC	-I- PTM: Phosphorylated on Ser-112 in response to survival stimuli.			
CC	Subsequent phosphorylation on Ser-136 promotes heterodimerization			
CC	with 14-3-3 proteins. This interaction then facilitates the			
CC	phosphorylation at Ser-155, a site within the BH3 domain, leading			
CC	to the release of Bcl-x(L) and the promotion of cell survival.			

ALIGNMENTS

Q9uix4 homo sapien
Q03030 salmonella
P13187 klebsiella
Q13049 homo sapien
P05886 simian immu
Q18823 caenorhabdi
Q9p0j0 h nadh-ubiq
P33602 escherichia
P33900 salmonella
O54918 mus musculu
O88498 rattus norv
P27757 simian immu

34 45 31.0 513 1 KCG1_HUMAN
35 45 31.0 590 1 DCOA_SALTY
36 45 31.0 595 1 DCOA_KLEPN
37 45 31.0 653 1 HT2A_HUMAN
38 45 31.0 865 1 ENV_SIVAT
39 45 31.0 1535 1 LML1_CAEEL
40 44.5 30.7 143 1 NB6M_HUMAN
41 44.5 30.7 907 1 NUOG_ECOLI
42 44.5 30.7 907 1 NUOG_SALTY
43 44 30.3 196 1 BIM_MOUSE
44 44 30.3 196 1 BIM_RAT
45 44 30.3 768 1 ENV_SIVAT

CC Ser-136 is the major site of AKT/PKB phosphorylation, Ser-155 the
 CC major site of protein kinase A (CAK) phosphorylation.
 CC -!- SIMILARITY: CONTAINS 1 BCL-2 homology 3 (BH3) domain.
 CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -----
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 CC -----
 DR EMBL; L37296; AAA64465.1; -
 DR PIR; A56671; A56671.
 DR HSSP; Q92934; IG571.
 DR MGD; MGI:1096330; Bad.
 DR InterPro; IPR000712; Bcl2 BH.
 DR PROSITE; PS01259; BH3; FALSE_NEG.
 KW Apoptosis; Phosphorylation.
 FT DOMAIN 147 161 BH3.
 FT MOD_RES 112 112 PHOSPHORYLATION (BY PKA AND PKB).
 FT MOD_RES 136 136 PHOSPHORYLATION (BY PKA AND PKB).
 FT MOD_RES 155 155 PHOSPHORYLATION (BY PKA AND PKB).
 FT MUTAGEN 112 112 S->A: NO PHOSPHORYLATION.
 FT MUTAGEN 136 136 S->A: NO PHOSPHORYLATION.
 FT MUTAGEN 155 155 S->A: NO PHOSPHORYLATION; INTERACTS WITH
 FT BCL-X(L).
 FT SQ SEQUENCE 204 AA; 22080 MW; 6C2BA910205053F7 CRC64;
 Query Match 100.0%; Score 145; DB 1; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.4e-14;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 NLWAAQRYGRLRMSDEFGSGFKGLP 27
 Db 140 NLWAAQRYGRLRMSDEFGSGFKGLP 166
 |||||
 RESULT 2
 ID BAD_RAT STANDARD; PRT; 205 AA.
 AC Q35147; Q70256; Q9JHX1.
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component
 DE 6) (Bcl-XL/Bcl-2 associated death promoter).
 GN BAD.
 OS Rattus norvegicus (Rat).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
 OX NCBI_Taxid=10116;
 RN [1]
 RP SEQUENCE FROM N.A., AND MUTAGENESIS OF SER-113 AND SER-137.
 RC TISSUE=Ovary;
 RX MEDLINE=98034386; PubMed=9369453; 1
 RA Hsu S.-Y., Kaipia A., Zhu L., Hsueh A.J.W.;
 RT "Interference of BAD (Bcl-XL/Bcl-2-associated death promoter)-induced
 RT apoptosis in mammalian cells by 14-3-3 isoforms and p11.";
 RL Mol. Endocrinol. 11:1858-1867(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=98194755; PubMed=9535132;
 RA D'Agata V., Magro G., Travali S., Musco S., Cavallaro S.;
 RT "Cloning and expression of the programmed cell death regulator BAD in
 RT the rat brain.";
 RL Neurosci. Lett. 243:137-140(1998).
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORMS ALPHA AND BETA).
 RC TISSUE=Brain;
 RX MEDLINE=21109372; PubMed=11161472;

RA Hammer S., Arumae U., Yu L.-Y., Sun Y.-P., Saarma M., Lindholm D.;
 RT "Functional characterization of two splice variants of rat BAD and
 RT their interaction with Bcl-w in sympathetic neurons.";
 RL Mol. Cell. Neurosci. 17:97-106(2001).
 CC -!- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 CC similarity). Appears to act as a link between growth factor
 CC receptor signaling and the apoptotic pathways.
 CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10. The Ser-
 CC 113/Ser-137 phosphorylated form binds 14-3-3 proteins.
 CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm (By similarity).
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing; Named isoforms=2;
 CC Name=Alpha;
 CC IsoId=Q35147-1; Sequence=Displayed;
 CC Name=Beta;
 CC IsoId=Q35147-2; Sequence=VSP_000534;
 CC -!- TISSUE SPECIFICITY: Expressed in all tissues tested, including
 CC brain, liver, spleen and heart. In the brain, restricted to
 CC epithelial cells of the choroid plexus. Isoform alpha is the more
 CC abundant form.
 CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -!- PTM: Phosphorylated on Ser-113 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-137 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-156, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-137 is the major site of AKT/PKB phosphorylation, Ser-156 the
 CC major site of protein kinase A (CAK) phosphorylation (By
 CC similarity).
 CC -!- SIMILARITY: CONTAINS 1 Bcl-2 homology 3 (BH3) domain.
 CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -----
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 CC -----
 DR EMBL; AF003523; AAC53374.1; -
 DR EMBL; AF031227; AAC15100.1; -
 DR EMBL; AF279910; AAF91427.1; -
 DR EMBL; AF279911; AAF91428.1; -
 DR HSSP; Q92934; IG571.
 DR InterPro; IPR000712; Bcl2 BH.
 DR PROSITE; PS01259; BH3; FALSE_NEG.
 KW Apoptosis; Phosphorylation; Alternative splicing.
 FT DOMAIN 148 162 BH3.
 FT MOD_RES 113 113 PHOSPHORYLATION (BY PKA AND PKB)
 FT MOD_RES 137 137 (BY SIMILARITY).
 FT MOD_RES 156 156 PHOSPHORYLATION (BY PKA AND PKB)
 FT MOD_RES 156 156 (BY SIMILARITY).
 FT VARSPLIC 166 205 LPRPKSAGTATCMROSASWTRIGSWDRNLKGSGTSPQ
 FT -> EELTYSVEFLPYRAIMEGNPLLSQSPSHLPPTTP
 FT /FTID=VSP_000534.
 FT MUTAGEN 113 113 S->A: NO EFFECT ON HETERODIMERIZATION
 FT WITH 14-3-3 PROTEINS.
 FT MUTAGEN 137 137 S->A: NO HETERODIMERIZATION WITH 14-3-3
 FT PROTEINS. NO EFFECT ON HETERODIMERIZATION
 FT WITH BCL2 NOR WITH PROTEIN P11.
 FT SDAGGR -> ERRGRK (IN REF. 1).
 FT CONFLICT 29 34 7AFA71DAE9CFA81 CRC64;
 FT SEQUENCE 205 AA; 22228 MW; 7AFA71DAE9CFA81 CRC64;

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Query Match      100.0%; Score 145; DB 1; Length 205;
Best Local Similarity 100.0%; Pred. No. 1.4e-14;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 NLWAAQRYGRLRMSDEFGSKGLP 27
Db 141 NLWAAQRYGRLRMSDEFGSKGLP 167

RESULT 3
BAD_HUMAN
ID BAD_HUMAN STANDARD; PRT; 168 AA.
AC Q92934; O14803;
DT 01-NOV-1997 (Rel. 35, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-
DE XL/Bcl-2 associated death promoter) (BCL2-like 8 protein).
GN BAD OR BBC6 OR BCL2L8.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Yin D.X., Li Z., Huang B., Chen S., Zhou H.;
RT "A human protein that interacts with Bcl-2 and have homology to mouse
RT BAD.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.
RX MEDLINE=97083574; PubMed=8929532;
RA Wang H.-G., Rapp U.R., Reed J.C.;
RT "Bcl-2 targets the protein kinase Raf-1 to mitochondria.";
RL Cell 87:629-638(1996).
RN [3]
RP SEQUENCE FROM N.A.
RA Takayama S., Reed J.C.;
RL Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A., AND DIMERIZATION.
RP TISSUE=Bone marrow;
RX MEDLINE=98049554; PubMed=9388232;
RA Ohtsuka S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,
RT "Dimerization properties of human BAD.";
RL J. Biol. Chem. 272:30866-30872(1997).
RN [5]
RP SEQUENCE FROM N.A.
RP TISSUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Haieff F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usslin T.B., Toshiyuki S., Carninci P., Frange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaby S.J.,
RA Rosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettner M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [6]

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RP STRUCTURE BY NMR OF 103-127.
RX MEDLINE=21073561; PubMed=11206074;
RA Petros A.M., Nettlesham D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
RA Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
RA Fesik S.W.;
RT "Rationale for Bcl-xL/Bad peptide complex formation from structure,
RT mutagenesis, and biophysical studies.";
RL Protein Sci. 9:2528-2534(2000).
CC -I- FUNCTION: Promotes cell death. Successfully competes for the
CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
CC of heterodimerization of these proteins with BAX. Can reverse the
CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
CC similarity). Appears to act as a link between growth factor
CC receptor signaling and the apoptotic pathways.
CC -I- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
CC The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
CC similarity).
CC -I- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
CC phosphorylation, locates to the cytoplasm.
CC -I- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
CC -I- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
CC BAX for their pro-apoptotic activity and for their interaction
CC with anti-apoptotic members of the Bcl-2 family.
CC -I- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
CC Subsequent phosphorylation on Ser-99 promotes heterodimerization
CC with 14-3-3 proteins. This interaction then facilitates the
CC phosphorylation at Ser-118, a site within the BH3 domain, leading
CC to the release of Bcl-x(L) and the promotion of cell survival.
CC Ser-99 is the major site of AKG/PKB phosphorylation, Ser-118 the
CC major site of protein kinase A (CAPK) phosphorylation (by
CC similarity).
CC -I- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC -I- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
CC -I- CAUTION: Ref.1 sequence differs from that shown due to frameshifts
CC in position 64 and 91.
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CC -----
DR EMBL; U66879; AAB36516.1; ALT_FRAME.
DR EMBL; AF021792; AAB72092.1; -.
DR EMBL; AF031523; AAB88124.1; -.
DR EMBL; BC001901; AAB01901.1; -.
DR PDB; 1G5J; 07-FEB-01.
DR Genbank; HGNC:936; BAD.
DR MIM; 603167; -.
DR GO; GO:0005737; Cytoplasm; NAS.
DR GO; GO:0005741; C-mitochondrial outer membrane; NAS.
DR GO; GO:0005515; F-protein binding activity; NAS.
DR GO; GO:0008632; P-apoptotic program; PAs.
DR GO; GO:0006917; P-induction of apoptosis; NAS.
DR InterPro; IPR000712; BCL2_BH.
DR PROSITE; PS01259; BH3; FALSE_NEG.
KW Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
FT DOMAIN 110 124
FT MOD_RES 75 75
FT PHOSPHORYLATION (BY PKA AND PKB) (BY
FT MOD_RES 99 99
FT PHOSPHORYLATION (BY PKA AND PKB) (BY
FT MOD_RES 118 118
FT PHOSPHORYLATION (BY PKA AND PKB) (BY
FT VARIANT 107 107
FT A->S (in dbSNP:3729933).
FT HELIX 106 121
FT SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
Query Match      83.1%; Score 120.5; DB 1; Length 168;

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SMART; SMO0327; VWFA; 1.
DR PROSITE; PS0234; VWFA; 1.
KW Serine protease inhibitor; Repeat; Signal; Multigene family;
FT GLYCOPROTEIN. 1 18 POTENTIAL.
FF SIGNAL 19 54 BY SIMILARITY.
FT CHAIN 55 702 INTER-ALPHA-TRYPSIN INHIBITOR HEAVY CHAIN H2.

PROPEP 303 946 BY SIMILARITY.
FF DOMAIN 308 468 VWFA.
FT CARBOHYD 118 118 N-LINKED (GLCNAC. .) (POTENTIAL).
FF CARBOHYD 263 263 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 445 445 N-LINKED (GLCNAC. .) (POTENTIAL).
FF CARBOHYD 578 578 N-LINKED (GLCNAC. .) (POTENTIAL).
FT BINDING 702 702 CHONDROITIN 4-SULFATE, CROSS-LINK SITE (BY SIMILARITY).

CONFLICT 510 510 V -> Y (IN REF. 2).
ET CONFLICT 595 595 E -> I (IN REF. 2).
FF SEQUENCE 946 AA; 106580 MW; CABEF5654587BZE CRC64;

Query Match 42.1%; Score 61; DB 1; Length 946;
Best Local Similarity 37.0%; Pred. No. 0.34;
Matches 10; Conservative 5; Mismatches 12; Indels 0; Gaps

QY 1 NLWAAQRYGRELARMDSDFEGSFKLP 27
:-| : | : ||| :-|:
DB 212 NWVIVELQGMRFLHVDPDTFEGHQVP 238

RESULT 5
ITH2_MOUSE STANDARD; PRT; 946 AA.
ID ITH2_MOUSE AC Q61703;
DC 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Inter-alpha-trypsin inhibitor heavy chain H2 precursor (ITI heavy chain H2) (inter-alpha-inhibitor heavy chain 2).
GN ITIH2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_Taxid=10090;
RN [1]
RC SEQUENCE FROM N.A.
RS STRAIN=C57BL/6N; TISSUE=Liver;
RX MEDLINE=95194326; PubMed=7534067;
RA Chan P., Risler J.-L., Raguenes G., Salier J.-P.;
RT "The three heavy-chain precursors for the inter-alpha-inhibitor family in mouse: new members of the multicopper oxidase protein group with differential transcription in liver and brain.";
RL Biochem. J. 306:505-512(1995).
CC !- FUNCTION: MAY ACT AS A CARRIER OF HYALURONAN IN SERUM OR AS A BINDING PROTEIN BETWEEN HYALURONAN AND OTHER MATRIX PROTEIN, INCLUDING THOSE ON CELL SURFACES IN TISSUES TO REGULATE THE LOCALIZATION, SYNTHESIS AND DEGRADATION OF HYALURONAN WHICH ARE ESSENTIAL TO CELLS UNDERGOING BIOLOGICAL PROCESSES (BY SIMILARITY).
CC !- SUBUNIT: I-ALPHA-I PLASMA PROTEASE INHIBITORS ARE ASSEMBLED FROM ONE OR TWO HEAVY CHAINS (H1, H2 OR H3) AND ONE LIGHT CHAIN, BIKUNIN. INTER-ALPHA-INHIBITOR (I-ALPHA-I) IS COMPOSED OF H1, H2 AND BIKUNIN. INTER-ALPHA-LIKE INHIBITOR (I'-ALPHA-LI) OF H2 AND BIKUNIN. AND PR-ALPHA-INHIBITOR (P-ALPHA-I) OF H3 AND BIKUNIN.
CC !- TISSUE SPECIFICITY: EXPRESSED IN BOTH LIVER AND BRAIN.
CC !- PTM: HEAVY CHAINS ARE INTERLINKED WITH BIKUNIN VIA A CHONDROITIN 4-SULFATE BRIDGE TO THEIR C-TERMINAL ASPARTATE (BY SIMILARITY).
CC !- SIMILARITY: BELONGS TO THE ITIH FAMILY.
CC !- SIMILARITY: Contains 1 VWFA domain.

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DR HSP: P00886; 1QK7.
DR SD: S000453; ARO4.
DR GO: GO:0003849; F:2-dehydro-3-deoxyphosphoheptonate aldolase . . . ; IDA.
DR InterPro: IPR006219; AROFGH.
DR InterPro: IPR006218; DAHP1/KDSA.
DR Pfam: PF00793; DAHP synth_1; 1.
DR ProDom: PD005060; AROFGH; 1.
DR TIGRfams: TIGR00034; AROFGH; 1.
KW Aromatic amino acid biosynthesis; Lyase; Multigene family.
SQ SEQUENCE 370 AA; 39749 MW; 594ED48F24175979 CRC64;

Query Match 35.98; Score 52; DB 1; Length 370;
Best Local Similarity 47.68; Pred. No. 2.7;
Matches 10; Conservative 5; Mismatches 6; Indels 0; Gaps 0;

Cy 1 NLWAAQRYGRELRLMSDFEG 21
   : ||| | : ||| |
Db 80 DLEAAQEAVALRLKLSDELKG 100

RESULT 8
ID AP3_ARATH STANDARD; PRT; 232 AA.
AC P35632; Q39003; Q8L879; Q9S703; Q9S014; Q9S015; Q9SQ16; Q9SQ17;
AC Q9S018; Q9S019; Q9S020; Q9S021; Q9S022; Q9SX13;
DI 01-JUN-1994 (Rel. 28, created)
DT 01-JUN-1994 (Rel. 29, last sequence update)
DT 15-SEP-2003 (Rel. 42, last annotation update)
DE Floral homeotic protein APETALA3.
GN AP3 OR AP3G54340 OR TL2E18.30.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Petal;
RX MEDLINE=92154682; PubMed=1346756;
RA Jack T., Brockman L.L., Meyerowitz E.M.;
RT "The homeotic gene APETALA3 of Arabidopsis thaliana encodes a MADS
RT box and is expressed in petals and stamens.";
RL Cell 58:583-597(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Landsberg erecta;
RX MEDLINE=95036018; PubMed=7948893;
RA Okamoto H., Yano A., Shiraiishi H., Okada K., Shimura Y.;
RT "Genetic complementation of a floral homeotic mutation, apetal3,
RT with an Arabidopsis thaliana gene homologous to DEFICIENS of
RT Antirrhinum majus.";
RL Plant Mol. Biol. 26:465-472(1994).
RN [3]
RP SEQUENCE FROM N.A., AND VARIANTS.
RC STRAIN=cv. Bla-1, cv. Bretagny, cv. Bs-1, cv. Bu-0, cv. Bu-2,
RC cv. Chi-1, cv. Co-1, cv. Columbia, cv. Corsacalla-1, cv. Cvi-0,
RC cv. Gr-3, cv. J1-1, cv. Kas-1, cv. Kent, cv. Landsberg erecta,
RC cv. Li-3, cv. Li-8, and cv. Lisse;
RX MEDLINE=99126449; PubMed=9927474;
RA Purugganan M.D., Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci: departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
RT genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RX MEDLINE=21016720; PubMed=11130713;
RA Salanoubat M., Lemcke K., Rieger M., Ansoorge W., Unseld M.,
RA Partmann B., Valle G., Bloecker H., Perez-Alonso M., Obermaier B.,
RA Delseny M., Boutry M., Griwell L.A., Mache R., Puigdomenech P.,
RA De Simone V., Choisine N., Artiguenave F., Robert C., Brottier P.,
RA Wincker P., Cattolico L., Weissenbach J., Saurin W., Quetier F.,

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RA Schaefer M., Mueller-Auer S., Gabel C., Fuchs M., Benes V.,
RA Wumbach E., Drzonek H., Erfle H., Jordan N., Bangert S.,
RA Wiedemann R., Kranz H., Voss H., Holland K., Brandt P., Yakatara G.,
RA Vezzi A., D'Angelo M., Pallavicini A., Toppo S., Simonati B.,
RA Conrad A., Hornischer K., Kauer G., Loehner T.-H., Nordsiek G.,
RA Reichelt J., Scharfe M., Schoen O., Bargues M., Terol J., Climent J.,
RA Navarro P., Collado C., Perez-Perez A., Ottenwaelder B., Duchemin D.,
RA Cooke R., Laudie M., Berger-Lauro C., Purnelle B., Masuy D.,
RA de Haan M., Maarse A.C., Alcaraz J.-P., Cottet A., Casacuberta E.,
RA Monfort A., Argirou A., Flores M., Liguori R., Vitale D.,
RA Manthaupt G., Haase D., Schoof H., Rüd S., Zaccaria P., Mewes H.-W.,
RA Mayer K.F.X., Kaul S., Town C.D., Koo H.L., Tallon L.J., Jenkins J.,
RA Rooney T., Rizzo M., Walts A., Utterback T., Fujii C.Y., Shea T.P.,
RA Creasy T.H., Haas B., Waiti R., Wu D., Peterson J., Van Aken S.,
RA Pal G., Miltscher J., Sellers P., Gill J.E., Feldblyum T.V.,
RA Preuss D., Lin X., Nierman W.C., Salzberg S.B., White O., Venter J.C.,
RA Fraser C.M., Kaneko T., Nakamura Y., Sato S., Kato T., Asamizu E.,
RA Sasamoto S., Kimura T., Idesawa K., Kawashima K., Kishida Y.,
RA Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
RA Nakayama S., Nakazaki N., Shinpo S., Takeuchi C., Wada T.,
RA Watanabe A., Yamada M., Yasuda M., Tabata S.;
RT "Sequence and analysis of chromosome 3 of the plant Arabidopsis
RT thaliana.";
RL Nature 408:820-822(2000).
RN [5]
RP SEQUENCE FROM N.A.
RC Brover V., Troukhan M., Alexandrov N., Lu Y.-P., Flavell R.,
RA Feldmann K.;
RT "Full-length cDNA from Arabidopsis thaliana.";
RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
RN [6]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RA Shinozaki K., Davis R.W., Ecker J.R., Theologis A.;
RT "RIKEN Arabidopsis full length cDNA clones (RAFLs) sequenced by the
RT SSP consortium (Salk/Stanford/PGECC).";
RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
RN [7]
RP SEQUENCE OF 36-128 FROM N.A.
RC STRAIN=cv. Columbia;
RX MEDLINE=99311297; PubMed=10382288;
RA Brunel D., Proger N., Pelletier G.;
RT "Development of amplified consensus genetic markers (ACGM) in Brassica
RT napus from Arabidopsis thaliana sequences of known biological
RT function.";
RL Genome 42:387-402(1999).
RN [8]
RP FUNCTION.
RX PubMed=8565821;
RA Krizek B.A., Meyerowitz E.M.;
RT "The Arabidopsis homeotic genes APETALA3 and PISTILLATA are sufficient
RT to provide the B class organ identity function.";
RL Development 122:11-22(1996).
RN [9]
RP CHARACTERIZATION.
RX PubMed=8643482;
RA Riechmann J.B., Krizek B.A., Meyerowitz E.M.;
RT "Dimerization specificity of Arabidopsis MADS domain homeotic proteins
RT APETALA1, APETALA3, PISTILLATA, and AGAMOUS.";
RL Proc. Natl. Acad. Sci. U.S.A. 93:4793-4798(1996).
RN [10]
RP GENETIC REGULATION.
RX PubMed=11283333;
RA Ng M., Yanofsky M.F.;
RT "Activation of the Arabidopsis B class homeotic genes by APETALA1.";
RL Plant Cell 13:739-753(2001).
RN [11]
RP CHARACTERIZATION.
RX PubMed=11206550;
RA Honma T., Goto K.;
RT "Complexes of MADS-box proteins are sufficient to convert leaves into
RT floral organs.";
RL Nature 409:525-529(2001).

```


-!- FUNCTION: Probable transcription factor involved in the genetic control of flower development.. Is required for normal development of petals and stamens in the wild-type flower. Forms an heterodimer with PISTILLATA that is required for autoregulation of both AP3 and PI genes. AP3/PI heterodimer interacts with APETAL3 or SEPALLATA3 to form a ternary complex that could be responsible for the regulation of the genes involved in the flower development.

-!- SUBUNIT: Forms an heterodimer with PISTILLATA, capable of binding to CARB-box sequences. AP3/PI heterodimer binds AP1 or SEP3 to form complexes.

-!- SUBCELLULAR LOCATION: Nuclear.

-!- TISSUE SPECIFICITY: Expressed in petals and stamens.

-!- INDUCTION: Positively regulated by the meristem identity proteins APETAL3 and LEAFY with the cooperation of UFO.

-!- MISCELLANEOUS: Mutations in AP3 cause transformation of petals into sepals and stamens into carpels.

-!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION FACTORS.

-!- SIMILARITY: Contains 1 K-box dimerization domain.

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EMBL; M86357; AAA32740.1; -
 DR EMBL; D21125; BAA04565.1; -
 DR EMBL; AF115798; AAD51867.1; -
 DR EMBL; AF115799; AAD51888.1; -
 DR EMBL; AF115800; AAD51889.1; -
 DR EMBL; AF115801; AAD51890.1; -
 DR EMBL; AF115802; AAD51891.1; -
 DR EMBL; AF115803; AAD51892.1; -
 DR EMBL; AF115804; AAD51893.1; -
 DR EMBL; AF115805; AAD51894.1; -
 DR EMBL; AF115806; AAD51895.1; -
 DR EMBL; AF115807; AAD51896.1; -
 DR EMBL; AF115808; AAD51897.1; -
 DR EMBL; AF115809; AAD51898.1; -
 DR EMBL; AF115810; AAD51899.1; -
 DR EMBL; AF115811; AAD51900.1; -
 DR EMBL; AF115812; AAD51901.1; -
 DR EMBL; AF115813; AAD51902.1; -
 DR EMBL; AF115814; AAD51903.1; -
 DR EMBL; AL132971; CAB81799.1; -
 DR EMBL; AY087369; RAM64919.1; -
 DR EMBL; AY070397; RAL49893.1; -
 DR EMBL; AV142590; RAN13159.1; -
 DR EMBL; AF056541; RAD41557.1; -
 DR PIR; A42095; A42095.
 DR HSSP; P11746; IMNM.
 DR TRANSFAC; T01776; -
 DR InterPro; IPR002487; TF_Kbox.
 DR InterPro; IPR002100; TF_MADSbox.
 DR Pfam; PF01486; K-box; 1.
 DR Pfam; PF00319; SRP-TF; 1.
 DR PRINTS; PR00404; MADSDOMAIN.
 DR SMART; SM00432; MADS; 1.
 DR PROSITE; PS00350; MADS_BOX_1; 1.
 DR PROSITE; PS00066; MADS_BOX_2; 1.
 KW Flowering; transcription regulation; Activator; Developmental protein;
 KW Nuclear protein; DNA-binding; Coiled coil; Polymorphism.
 FT DOMAIN 3 57 MADS.
 FT DOMAIN 93 165 K-BOX.
 FT DOMAIN 75 164 COILED COIL (POTENTIAL).
 FT VARIANT 31 31 K -> R (in strain cv. Lisse).
 FT VARIANT 47 47 M -> T (in strain cv. Bretagne).
 FT VARIANT 61 61 N -> D (in strain cv. Corsacalla-1).
 FT VARIANT 73 73 T -> S (in strain cv. Li-8).

FT VARIANT 109 109 L -> V (in strain cv. Kas-1).
 FT VARIANT 115 115 E -> K (in strains cv. Chl-1 and cv. Gr-3).
 FT

Query Match 35.2%; Score 51; DB 1; Length 232;
 Best Local Similarity 44.4%; Pred. No. 2.3;
 Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRMSDEFGSPK 24
 ||| :|||: ||| :||
 DB 107 QRLGCLDELDIQELRRLEDEMENTFK 133

RESULT 9
 CE05_MOUSE
 ID CE05_MOUSE STANDARD; PRT; 851 AA.
 AC Q8K2H3;
 DT 15-SEP-2003 (Rel. 42, Created)
 DT 15-SEP-2003 (Rel. 42, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Protein C5orf5 homolog.
 GN C5ORF5.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=FVB/N;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins K.F., Jordan H., Moore T., Max S.I., Wang J., Heieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Roderfeld Y.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 CC -!- SIMILARITY: Belongs to the FAM13 family.
 CC -!- SIMILARITY: Contains 1 Rho-GAP domain.
 CC

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EMBL; BC031465; AAH31465.1; -
 DR InterPro; IPR000198; RhoGAP.
 DR Pfam; PF00620; RhoGAP; 1.
 DR SMART; SM00324; RHOAP; 1.
 DR PROSITE; PS0238; RHOAP; 1.
 KW GTPase activation.
 FT DOMAIN 23 212 RHO-GAP.
 FT DOMAIN 189 256 GLU-RICH.
 SQ SEQUENCE 851 AA; 97054 MW; C2B26669FB6DB2CE CRC64;

Query Match 34.5%; Score 50; DB 1; Length 851;
 Best Local Similarity 45.5%; Pred. No. 13;

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Matches 10; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 2 LWAAQYGRRLRMSDEFGSF 23
   ||| :||| :||| :||| :
Db 784 LWKARAEKKLRMLRFEFEAF 805

RESULT 10
MATK_LEDPA STANDARD; PRT; 506 AA.
AC O62992;
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Maturase K (intron maturase).
GN MATK.
OS Ledum palustre (Wild rosemary).
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75583;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of sectional relationships in the genus
RT Rhododendron (Ericaceae) based on matk sequences.";
RL Shokubutsu Kenkyu Zasshi 73:143-154(1998).
CC -!- FUNCTION: Probably assists in splicing chloroplast group II
CC introns (By similarity).
CC -!- SIMILARITY: BELONGS TO THE INTRON MATURASE FAMILY 2. MATK
CC SUBFAMILY.
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CC -----
CC EMBL; AB012741; BAA25862.1; -
CC InterPro; IPR000442; Intron_maturase2.
CC InterPro; IPR002866; MatK_N.
CC Pfam; PF01348; Intron_maturase2; 1.
CC Pfam; PF01824; MatK_N; 1.
CC mRNA processing; Chloroplast.
CC KW Maturase K (intron maturase).
CC ADAA4325E92436B8 CRC64;
CC SEQUENCE 506 AA; 60412 MW; CFEA926307DAC85E CRC64;
CC -----
Query Match 34.1%; Score 49.5; DB 1; Length 506;
Best Local Similarity 36.7%; Pred. No. 9;
Matches 11; Conservative 5; Mismatches 7; Indels 7; Gaps 1;

QY 2 LWAA-----QRYGRLRMSDEFGSKF 24
   |||| :||| :||| :||| :
Db 393 VWAALSDSDIERFGRYRNLSHYSGSLK 422

RESULT 11
MATK_RHOFR STANDARD; PRT; 506 AA.
AC O62984;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Maturase K (intron maturase).
GN MATK.
OS Rhododendron ferrugineum (Alpenrose).
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49629;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of sectional relationships in the genus
RT Rhododendron (Ericaceae) based on matk sequences.";
RL Shokubutsu Kenkyu Zasshi 73:143-154(1998).
CC -!- FUNCTION: Probably assists in splicing chloroplast group II
CC introns (By similarity).
CC -!- SIMILARITY: BELONGS TO THE INTRON MATURASE FAMILY 2. MATK
CC SUBFAMILY.
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CC -----
CC EMBL; AB012751; BAA25872.1; -
CC InterPro; IPR000442; Intron_maturase2.
CC InterPro; IPR002866; MatK_N.
CC Pfam; PF01348; Intron_maturase2; 1.
CC Pfam; PF01824; MatK_N; 1.
CC mRNA processing; Chloroplast.
CC KW Maturase K (intron maturase).
CC ADAA4325E92436B8 CRC64;
CC SEQUENCE 506 AA; 60412 MW; CFEA926307DAC85E CRC64;
CC -----
Query Match 34.1%; Score 49.5; DB 1; Length 506;
Best Local Similarity 36.7%; Pred. No. 9;
Matches 11; Conservative 5; Mismatches 7; Indels 7; Gaps 1;

QY 2 LWAA-----QRYGRLRMSDEFGSKF 24
   |||| :||| :||| :||| :
Db 393 VWAALSDSDIERFGRYRNLSHYSGSLK 422

RESULT 12
MATK_RHOTS STANDARD; PRT; 506 AA.
AC O62991;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Maturase K (intron maturase).
GN MATK.
OS Rhododendron tsusiiophyllum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49629;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of sectional relationships in the genus
RT Rhododendron (Ericaceae) based on matk sequences.";
RL Shokubutsu Kenkyu Zasshi 73:143-154(1998).
CC -!- FUNCTION: Probably assists in splicing chloroplast group II
CC introns (By similarity).
CC -!- SIMILARITY: BELONGS TO THE INTRON MATURASE FAMILY 2. MATK
CC SUBFAMILY.
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CC -----
CC EMBL; AB012741; BAA25862.1; -
CC InterPro; IPR000442; Intron_maturase2.
CC InterPro; IPR002866; MatK_N.
CC Pfam; PF01348; Intron_maturase2; 1.
CC Pfam; PF01824; MatK_N; 1.
CC mRNA processing; Chloroplast.
CC KW Maturase K (intron maturase).
CC ADAA4325E92436B8 CRC64;
CC SEQUENCE 506 AA; 60534 MW; ADAA4325E92436B8 CRC64;
CC -----
Query Match 34.1%; Score 49.5; DB 1; Length 506;
Best Local Similarity 36.7%; Pred. No. 9;
Matches 11; Conservative 5; Mismatches 7; Indels 7; Gaps 1;

QY 2 LWAA-----QRYGRLRMSDEFGSKF 24
   |||| :||| :||| :||| :
Db 393 VWAALSDSDIERFGRYRNLSHYSGSLK 422

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DR EMBL; AB012750; BAA25871.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01346; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60569 MW; ABE12FF6809C233E CRC64;

Query Match          34.1%; Score 49.5; DB 1; Length 506;
Best Local Similarity 36.7%; Pred. No. 9;
Matches 11; Conservative 5; Mismatches 7; Indels 7; Gaps 1;

QY 2 LWAA-----QRYGRLRMSDEFGSKF 24
   :||| :||| :||| :||| :|||
Db 393 VWAALSDSDIIERFGRIYNLHYSGSLK 422

RESULT 13
END8_ECO57
ID END8_ECO57 STANDARD; PRT; 262 AA.
AC Q8X9C6.
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Endonuclease VII (EC 3.2.-.-).
GN NEI OR Z0865 OR EC50739.
OS Escherichia coli O157:H7.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=83334;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / EDL933 / ATCC 700927;
RX MEDLINE=21074935; PubMed=11206551;
RA Perna N.T., Plunkett G., Burland V., Mau B., Glasner J.D.,
RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
RA Posfai G., Hackett J., Klink S., Boutin A., Shao Y., Miller I.,
RA Grobeck E.J., Davis N.W., Lim A., Dimalanta E.T., Potamouzis K.,
RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,
RA Welch R.A., Blattner F.R.;
RT "Genome sequence of enterohaemorrhagic Escherichia coli O157:H7.";
RL Nature 409:529-533(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / RMD 0509952;
RX MEDLINE=21156231; PubMed=11258796;
RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,
RA Iida T., Takami H., Honda T., Sasakawa C., Ogasawara N., Yasunaga T.,
RA Kuhara S., Shiba T., Hattori M., Shinagawa H.;
RT "Complete genome sequence of enterohaemorrhagic Escherichia coli
RL O157:H7 and genomic comparison with a laboratory strain K-12.";
EL DNA Res. 8:11-22(2001).
CC -1- FUNCTION: DNA N-GLYCOSYLASE WITH AN AP LYASE ACTIVITY. REQUIRED
CC FOR THE REPAIR OF OXIDATIVE DNA DAMAGE. CLEAVES THE DNA BACKBONE
CC BY BETA-DELTA ELIMINATION AS WELL AS 5'DEOXYRIBOSE PHOSPHATE (BY
CC SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE FPG FAMILY.
CC -----
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CC -----
DR EMBL; AE005249; AAG5037.1; -.
DR EMBL; AF002552; BAB34162.1; -.
DR PIR; A85572; A85572.
DR PIR; C90721; C90721.
DR InterPro; IPR000191; Fapy_DNA_glyco.
DR InterPro; IPR000214; Fapy_DNAGlyco_zn.

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DR Pfam; PF01149; Fapy_DNA_glyco; 1.
DR ProDom; PD003680; Fapy_DNA_glyco; 1.
DR PROSITE; PS01242; FPG; 1.
KW DNA repair; Hydrolase; Glycosidase; Endonuclease; Zinc; Zinc-finger;
KW Complete proteome.
FT INIT MET 0 BY SIMILARITY.
FT ZN_FING 237 260 POTENTIAL.
SQ SEQUENCE 262 AA; 29680 MW; 37C3C5E236E07A88 CRC64;

Query Match          33.8%; Score 49; DB 1; Length 262;
Best Local Similarity 47.4%; Pred. No. 5.2;
Matches 9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 9 GRELRMSDEFGSKFGLP 27
   | :||| :||| :|||
Db 3 GFEIRRAADNLEAAIKGP 21

RESULT 14
END8_ECOLI
ID END8_ECOLI STANDARD; PRT; 262 AA.
AC P50455;
DT 01-OCT-1996 (Rel. 34, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Endonuclease VIII (EC 3.2.-.-).
GN NEI OR B0714.
OS Escherichia coli.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=562;
RN [1]
RP SEQUENCE FROM N.A. AND SEQUENCE OF 1-35; 188-205 AND 213-226.
RX MEDLINE=97315255; PubMed=9171429;
RA Jiang D., Hatanet Z., Blaisdell J.O., Melamede R.J., Wallace S.S.;
RT "Escherichia coli endonuclease VIII: cloning, sequencing, and
RT overexpression of the nei structural gene and characterization of nei
RT and nei nth mutants.";
RL J. Bacteriol. 179:3773-3782(1997).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=97315256; PubMed=9171430;
RA Saito Y., Uraki F., Nakajima S., Asaeda A., Ono K., Kubo K.,
RA Yamamoto K.;
RT "Characterization of endonuclease III (nth) and endonuclease VIII
RT (nei) mutants of Escherichia coli K-12.";
RL J. Bacteriol. 179:3783-3785(1997).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blattner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12.";
RL Science 277:1453-1474(1997).
RN [4]
RP SEQUENCE FROM N.A.
RA Mori H.;
RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: DNA N-GLYCOSYLASE WITH AN AP LYASE ACTIVITY. REQUIRED
CC FOR THE REPAIR OF OXIDATIVE DNA DAMAGE. CLEAVES THE DNA BACKBONE
CC BY BETA-DELTA ELIMINATION AS WELL AS 5'DEOXYRIBOSE PHOSPHATE.
CC -1- SIMILARITY: BELONGS TO THE FPG FAMILY.
CC -----
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CC -----
DR EMBL; U38616; AAC45355.1; -.
DR EMBL; D89754; BAA20414.1; -.
DR EMBL; AE000174; AAC73808.1; -.
DR EMBL; D90710; BAA35378.1; -.
DR PIR; A64807; A64807.
DR PDB; 1K3W; 04-OCT-02.
DR Ecogene; EGI3237; nei.
DR InterPro; IPR000191; Fapy_DNA_glyco.
DR InterPro; IPR000214; Fapy_DNAGlyco_zn.
DR Pfam; PF01149; Fapy_DNA_glyco; 1.
DR ProDom; PD003680; Fapy_DNA_glyco; 1.
DR PROSITE; PS01242; FPG; 1.
DR DNA repair; Hydrolase; Glycosidase; Endonuclease; Zinc; Zinc-finger;
KW Complete proteome; 3d-structure.
FT INIT_MET 0
FT ZN_FING 237 260 POTENTIAL.
SQ SEQUENCE 262 AA; 29714 MW; 5010961768ADC265 CRC64;

Query Match 33.8%; Score 49; DB 1; Length 262;
Best Local Similarity 47.4%; Pred. No. 5.2;
Matches 9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 9 GRELRMSDEFECSFKGLP 27
   | | | | | | | | | |
Db 3 GPEIRREADNLEAIKGP 21

RESULT 15
END8_SALTI
ID END8_SALTI STANDARD; PRT; 262 AA.
AC Q828D2;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Endonuclease VIII (EC 3.2.-.-).
GN NEI OR STY0771 OR T2148.
OS Salmonella typhi.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID=601;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CT18;
RX MEDLINE=21534947; PubMed=11677608;
RA Parkhill J., Dougan G., James K.D., Thomson N.R., Pickard D., Wain J.,
RA Churchhill J., Mungall K.L., Bentley S.D., Holden M.T.G., Sebahia M.,
RA Baker S., Basham D., Brooks K., Chillingworth T., Connor P.,
RA Cronin A., Davis P., Davies R.M., Dowd L., White N., Farrar J.,
RA Feltwell T., Hamlin N., Haque A., Hien T.T., Holroyd S., Jagels K.,
RA Krogh A., Larsen T.S., Leather S., Moule S., O'Gaora P., Parry C.,
RA Quail M., Rutherford K., Simmonds M., Skelton J., Stevens K.,
RA Whitehead S., Barrell B.G.;
RT "Complete genome sequence of a multiple drug resistant Salmonella
RT enterica serovar Typhi CT18.";
RL Nature 413:848-852(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Ty2 / ATCC 700931;
RX MEDLINE=22531367; PubMed=12644504;
RA Deng W., Liou S.-R., Plunkett G. III, Mayhew G.F., Rose D.J.,
RA Burland V., Kodoyianni V., Schwartz D.C., Blattner F.R.;
RT "Comparative genomics of Salmonella enterica serovar Typhi strains Ty2
RT and CT18.";
RL J. Bacteriol. 185:2330-2337(2003).
CC -!- FUNCTION: DNA N-GLYCOYLASE WITH AN AP LYASE ACTIVITY. REQUIRED
CC FOR THE REPAIR OF OXIDATIVE DNA DAMAGE. CLEAVES THE DNA BACKBONE
CC BY BETA-DELTA ELIMINATION AS WELL AS 5'DEOXYRIBOSE PHOSPHATE (BY
CC similarity).
CC -!- SIMILARITY: BELONGS TO THE FPG FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; AL627268; CAD05190.1; -.
DR EMBL; AE016841; AAO89782.1; -.
DR InterPro; IPR000191; Fapy_DNA_glyco.
DR InterPro; IPR000214; Fapy_DNAGlyco_zn.
DR Pfam; PF01149; Fapy_DNA_glyco; 1.
DR ProDom; PD003680; Fapy_DNA_glyco; 1.
DR PROSITE; PS01242; FPG; 1.
DR DNA repair; Hydrolase; Glycosidase; Endonuclease; Zinc; Zinc-finger;
KW Complete proteome.
FT INIT_MET 0
FT ZN_FING 237 260 BY SIMILARITY.
FT ZN_FING 237 260 POTENTIAL.
SQ SEQUENCE 262 AA; 29734 MW; 4E255C0CCF59A6A3 CRC64;

Query Match 33.8%; Score 49; DB 1; Length 262;
Best Local Similarity 47.4%; Pred. No. 5.2;
Matches 9; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 9 GRELRMSDEFECSFKGLP 27
   | | | | | | | | | |
Db 3 GPEIRREADNLEAIKGP 21

Search completed: September 15, 2003, 17:22:59
Job time : 6.36429 secs
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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 ; Search time 29.3143 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-3

Perfect score: 145

Sequence: 1 NLWAAQRYGRELRRMSDFEGSKGLP 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SPTREMBL_23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phase:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_rvirus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	87	60.0	146	13	Q9I9N2
2	60	41.4	946	11	Q8K016
3	53	36.6	196	16	Q8VJS3
4	53	36.6	223	16	Q10843
5	52	35.9	471	17	Q8ZV71
6	52	35.9	946	6	Q9GLY6
7	51	35.2	231	10	Q9SEGO
8	51	35.2	232	10	Q9SQ20
9	51	35.2	232	10	Q9SQ22
10	51	35.2	232	10	Q9SQ17
11	51	35.2	232	10	Q9SQ19
12	51	35.2	232	10	Q9SQ21
13	51	35.2	232	10	Q8LB79
14	51	35.2	232	10	Q9SQ16
15	51	35.2	232	10	Q9SQ15
16	51	35.2	232	10	Q9S7Q3

17	51	35.2	232	10	Q9SQ18
18	50.5	34.8	904	2	Q9KGW3
19	50.5	34.8	909	16	Q8E134
20	50.5	34.8	990	10	Q9C793
21	50	34.5	168	11	Q8K316
22	50	34.5	283	15	Q37056
23	50	34.5	374	17	Q9HN29
24	50	34.5	516	10	Q9SP5
25	50	34.5	851	11	Q8K2H3
26	49.5	34.1	153	5	Q9UB33
27	49.5	34.1	401	5	Q97407
28	49.5	34.1	505	8	Q47148
29	49.5	34.1	506	8	Q47149
30	49.5	34.1	506	8	Q47171
31	49.5	34.1	506	8	Q63960
32	49.5	34.1	506	8	Q62982
33	49.5	34.1	506	8	Q62975
34	49.5	34.1	506	8	Q62972
35	49.5	34.1	506	8	Q62989
36	49.5	34.1	506	8	Q62978
37	49.5	34.1	506	8	Q47155
38	49.5	34.1	506	8	Q47152
39	49.5	34.1	506	8	Q47173
40	49.5	34.1	506	8	Q62990
41	49.5	34.1	506	8	Q62974
42	49.5	34.1	506	8	Q62993
43	49.5	34.1	506	8	Q47170
44	49.5	34.1	506	8	Q47174
45	49.5	34.1	506	8	Q62983

ALIGNMENTS

RESULT 1
Q9I9N2 PRELIMINARY; PRT; 146 AA.
AC Q9I9N2
DT 01-OCT-2000 (Tremblrel. 15, Created)
DT 01-DEC-2001 (Tremblrel. 19, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Bad.
GN BAD.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A. PubMed=10917738;
RX MEDLINE=20373792; PubMed=10917738;
RA Inohara N., Nunez G.;
RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish";
RL Cell Death Differ. 7:509-510(2000).
DR EMBL; AF231017; AAF66962.2;
DR HSP; Q22934; IG57.
DR ZFIN; ZDB-GENE-000616-1; bad.
SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;
Query Match 60.0%; Score 87; DB 13; Length 146;
Best Local Similarity 65.2%; Pred. No. 6.7e-05;
Matches 15; Conservative 5; Mismatches 3; Indels 0; Gaps 0;
QY 2 LWAQRYGRELRRMSDFEGSK 24
|||||:|||||:
Db 89 LWAQRYGRELRRMSDFEGSK 111
RESULT 2
Q8K016 PRELIMINARY; PRT; 946 AA.
ID Q8K016
AC Q8K016;


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Query Match      35.9%; Score 52; DB 17; Length 471;
Best Local Similarity 41.7%; Pred. No. 34;
Matches 10; Conservative 3; Mismatches 11; Indels 0; Gaps 0;

Qy 3 WAAQRYGRELRRMSDEFGSKGL 26
Db 404 WQHSGMGRELRLMRAETAEIAGEFGAL 427

RESULT 6
Q9GLY6 PRELIMINARY; PRT; 946 AA.
AC Q9GLY6;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Inter-alpha-trypsin inhibitor heavy chain2.
GN INT-HC2
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
RP SEQUENCE FROM N.A.
RA Suzuki A.;
RT "Rabbit inter-alpha-trypsin inhibitor heavy chain.";
RL Submitted (OCT-2000) to the EMBL/GenBank/DDJB databases.
DR EMBL; AB050592; BAB17301.1; -
DR InterPro; IPR006587; VIT.
DR InterPro; IPR002035; VWF_A.
DR Pfam; PF00092; vwa; 1
DR SMART; SM00609; VIT; 1.
DR SMART; SM00327; VWA; 1.
DR PROSITE; PS00234; VWFA; 1.
SQ SEQUENCE 946 AA; 106240 MW; B7AF05434B228CC5 CRC64;

Query Match      35.98%; Score 52; DB 6; Length 946;
Best Local Similarity 33.38%; Pred. No. 74;
Matches 9; Conservative 13; Mismatches 13; Indels 0; Gaps 0;

Qy 1 NLWAAQRYGRELRRMSDEFGSKGLF 27
Db 212 DWVILPQGLRFLHVPDFSGHFDGVP 238

RESULT 7
Q9SEGO PRELIMINARY; PRT; 231 AA.
AC Q9SEGO;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Apetala3 (Fragment).
OS Arabidopsis lyrata.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=59689;
RN [1]
RP SEQUENCE FROM N.A.
RA Layton-Rauh A.L.; Buckler E.S. IV, Purugganan M.D.;
RT "Patterns of molecular evolution among paralogous floral homeotic genes.";
RL Mol. Biol. Evol. 16:1037-1045(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION FACTORS.
DR EMBL; AF143380; AAF25590.1; -.
DR HSSP; P11746; IMNW.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.

Qy 6 QRYG-----RELRRMSDEFGSKF 24
Db 107 QRLGECGLDELIDQLRLEDEMENTFK 133

RESULT 9
Q9SQ22 PRELIMINARY; PRT; 232 AA.
AC Q9SQ22;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Floral homeotic protein AP3.

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DR Pfam; PF00319; SRF-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS0066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
FT NON_TER 231
SQ SEQUENCE 231 AA; 27176 MW; A67CAELEBBD8F7AA CRC64;

Query Match      35.2%; Score 51; DB 10; Length 231;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

Qy 6 QRYG-----RELRRMSDEFGSKF 24
Db 107 QRLGECGLDELIDQLRLEDEMENTFK 133

RESULT 8
Q9SQ20 PRELIMINARY; PRT; 232 AA.
AC Q9SQ20;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Corscalla;
RX MEDLINE=99126449; PubMed=9927474;
RA Purugganan M.D., Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures from the equilibrium-neutral model at the APETALA3 and PISTILLATA genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION FACTORS.
DR EMBL; AF115806; AAD51895.1; -.
DR HSSP; P11746; IMNW.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRF-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS0066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27342 MW; BDFDCB59B73F4601 CRC64;

Query Match      35.2%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

Qy 6 QRYG-----RELRRMSDEFGSKF 24
Db 107 QRLGECGLDELIDQLRLEDEMENTFK 133

RESULT 9
Q9SQ22 PRELIMINARY; PRT; 232 AA.
AC Q9SQ22;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Floral homeotic protein AP3.

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OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OX eurosids II; Brassicales; Brassicaceae; Arabidopsis.
RN NCBI_TaxID=3702;
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Kent;
RX MEDLINE=99126449; PubMed=9927474;
RA Purgananan M.D.; Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
RT genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AF115805; AAD51894.1; -.
DR HSP; P11746; 1NMN.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR PRINTS; PF00319; SRF-TF; 1.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27286 MW; 66976305B88B63E3 CRC64;

Query Match 35.2%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

Qy 6 QRYG-----RELRRMSDEFEFSK 24
Db 107 QRLGCLDELDIQELRRLEDEMENTFK 133

RESULT 13
Q8LB79 ID Q8LB79 PRELIMINARY; PRT; 232 AA.
AC Q8LB79;
DT 01-OCT-2002 (TREMBlrel. 22, Created)
DT 01-OCT-2002 (TREMBlrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Floral homeotic protein APETALA3 (AP3).
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN SEQUENCE FROM N.A.
RP SEQUENCE FROM N.A.
RA Haas B.J.; Volfovsky N.; Town C.D.; Troupkan M.; Alexandrov N.,
RA Feldmann K.A.; Flavell R.B.; White O.; Salzberg S.L.;
RT "Full-length messenger RNA sequences greatly improve genome
RT annotation.";
RL Genome Biol. 0:0-0(2002).
RN [2]
RP SEQUENCE FROM N.A.
RA Brover V.; Troupkan M.; Alexandrov N.; Lu Y.-P.; Flavell R.,
RA Feldmann K.;
RT "Full-length cDNA from Arabidopsis thaliana.";
RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AY087369; AM64919.1; -.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR PRINTS; PF00319; SRF-TF; 1.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27314 MW; DB8CAFC835557D6 CRC64;

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DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27339 MW; CC9703F959CFAD5 CRC64;

Query Match 35.2%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

Qy 6 QRYG-----RELRRMSDEFEFSK 24
Db 107 QRLGCLDELDIQELRRLEDEMENTFK 133

RESULT 14
Q9SQ16 ID Q9SQ16 PRELIMINARY; PRT; 232 AA.
AC Q9SQ16;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN SEQUENCE FROM N.A.
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Lisse;
RX MEDLINE=99126449; PubMed=9927474;
RA Purgananan M.D.; Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
RT genes of Arabidopsis thaliana.";
RL Genetics 151:839-848(1999).
CC -|- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -|- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AF115810; AAD51899.1; -.
DR HSP; P11746; 1NMN.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR PRINTS; PF00319; SRF-TF; 1.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS50066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
SQ SEQUENCE 232 AA; 27314 MW; DB8CAFC835557D6 CRC64;

Query Match 35.2%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

Qy 6 QRYG-----RELRRMSDEFEFSK 24
Db 107 QRLGCLDELDIQELRRLEDEMENTFK 133

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AC Q9SQ15;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Floral homeotic protein AP3.
GN APETALA3.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

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OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_taxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Kas-1;
RX MEDLINE=99126449; PubMed=9927474;
RA Purugganan M.D.; Suddith J.I.;
RT "Molecular population genetics of floral homeotic loci. Departures
RT from the equilibrium-neutral model at the APETALA3 and PISTILLATA
RT genes of Arabidopsis thaliana."
RL Genetics 151:839-848(1999).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
CC FACTORS.
DR EMBL; AF115812; AA051901.1; -.
DR HSSP; P11746; IMNM.
DR InterPro; IPR002487; TF_Kbox.
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DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRF-TF; 1.
DR PRINTS; PRO0404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
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DR PROSITE; PS00066; MADS_BOX_2; 1.
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
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Query Match 35.2%; Score 51; DB 10; Length 232;
Best Local Similarity 44.4%; Pred. No. 22;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 6 QRYG-----RELRLMSDEFECSFK 24
DB ||| :||||| :||
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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:01 ; Search time 39.6 Seconds
(without alignments)
112.231 Million cell updates/sec

Title: US-09-544-664-55

Perfect score: 148

Sequence: 1 KNLWAAQRYGRELRLMSDEFSGFKGLK/28

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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24: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	148	100.0	28	21 AAB37055	Bcl2 polypeptide B
2	143	96.6	27	21 AAB37056	Bcl2 polypeptide B
3	138	93.2	26	21 AAB37001	Bcl2 polypeptide B
4	138	93.2	26	21 AAB37002	Bcl2 polypeptide B
5	138	93.2	27	21 AAB37003	Bcl2 polypeptide B
6	138	93.2	162	22 AAB70370	Shorter murine BAD
7	138	93.2	204	17 AAR95168	Bcl-x(L)/bcl-2 ass
8	138	93.2	204	19 AAW61315	Murine BCL-XL/BCL-
9	138	93.2	204	19 AAW61316	Mutant BCL-XL/BCL-

10	138	93.2	204	19 AAW61317	Mutant BCL-XL/BCL-
11	138	93.2	204	19 AAW61318	Mutant BCL-XL/BCL-
12	138	93.2	204	19 AAW58832	Murine BAD protein
13	138	93.2	204	22 AAB70369	Longer murine BAD
14	138	93.2	204	24 ABR39082	Murine BAD protein
15	138	93.2	204	24 ABR39082	Bad-DTR apoptosis
16	114	77.0	24	23 AAU00220	Human Bad peptide
17	114	77.0	25	23 AAB56161	Human Bad peptide
18	114	77.0	25	23 AAB56161	PPC-interacting T
19	114	77.0	25	23 AAB56161	Mutant Bcl2 compet
20	114	77.0	25	23 AAB56161	Mutant Bcl2 compet
21	114	77.0	25	23 AAB56161	Human Bad peptide
22	114	77.0	25	23 AAB56161	Human Bad peptide
23	114	77.0	25	23 AAB56161	BBC6 protein, for r
24	114	77.0	25	23 AAB56161	Human Bcl-XL/Bcl-2
25	114	77.0	25	23 AAB56161	Human cell prolif
26	114	77.0	25	23 AAB56161	Human BAD mutant a
27	114	77.0	25	23 AAB56161	Human BAD protein.
28	114	77.0	25	23 AAB56161	Amino acid sequenc
29	114	77.0	25	23 AAB56161	Human BAD protein
30	113	76.4	23	17 AAR95166	Human ovarian anti
31	111	75.0	25	23 AAB78490	bcl-x(L)/bcl-2 ass
32	111	75.0	25	23 AAB78490	Mutant Bcl2 compet
33	110	74.3	25	23 AAB78488	Human Bad peptide
34	110	74.3	25	23 AAB78488	Mutant Bcl2 compet
35	110	74.3	25	23 AAB78488	Mutant Bcl2 compet
36	110	74.3	25	23 AAB78488	Human Bad peptide
37	109	73.6	23	23 AAU78616	Human Bad peptide
38	109	73.6	25	23 AAB78486	Human Bad peptide
39	109	73.6	25	23 AAB78486	Mutant Bcl2 compet
40	109	73.6	25	23 AAB78486	Mutant Bcl2 compet
41	109	73.6	25	23 AAB78486	Mutant Bcl2 compet
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43	109	73.6	25	23 AAB78486	Human Bad peptide
44	108	73.0	25	23 AAB78485	Mutant Bcl2 compet
45	108	73.0	25	23 AAB78485	Human Bad peptide

ALIGNMENTS

RESULT 1
AAB37055
ID AAB37055 standard; peptide; 28 AA.
XX AAB37055;
XX AC
XX AC
DT 28-FEB-2001 (first entry)
DE Bcl2 polypeptide BH3 domain peptide #55.
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.
XX Homo sapiens.
XX OS
XX WC2000059526-A1.
XX PD 12-OCT-2000.
XX PD 06-APR-2000; 2000WO-US09352.
XX PR 07-APR-1999; 99US-0128202.
XX PA (UYJB-) UNIV JEFFERSON THOMAS.
XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX WPI; 2000-679325/66.
XX

XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer
XX
XX
PS Claim 18; Page 19; 74pp; English.
XX
XX The invention relates to a peptide conjugate having the formula:
CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl optionally
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC of the peptide portion of the conjugate. The peptides represent analogues
CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
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Query Match 100.0%; Score 148; DB 21; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.4e-15;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 1 KNLWAAQRYGRELRRMSDEFGSKGLK 28
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XX
XX AAB37056;
AC
XX 28-FEB-2001 (first entry)
DT
XX Bcl2 polypeptide BH3 domain peptide #56.
DE
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.
XX
XX Homo sapiens.
OS
XX W0200059526-A1.
PN
XX 12-OCT-2000.
PD
XX 06-APR-2000; 2000WO-US09352.
PF
XX 07-APR-1999; 99US-0128202.
XX
XX (UYJE-) UNIV JEFFERSON THOMAS.
PA
XX

PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX WPI; 2000-679325/66.
XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer
XX
XX Claim 18; Page 19; 74pp; English.
XX
XX The invention relates to a peptide conjugate having the formula:
CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the peptide portion of the conjugate. The peptides represent analogues
CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
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|||||
DB 1 KNLWAAQRYGRELRRMSDEFGSKGL 27
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AAB37001
ID AAB37001 standard; peptide; 26 AA.
XX
XX AAB37001;
AC
XX 28-FEB-2001 (first entry)
DT
XX Bcl2 polypeptide BH3 domain peptide #1.
DE
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.
XX
XX Homo sapiens.
OS
XX W0200059526-A1.
PN
XX 12-OCT-2000.
PD
XX 06-APR-2000; 2000WO-US09352.
PF
XX 07-APR-1999; 99US-0128202.
XX
XX

XX PA (UYJE-) UNIV JEFFERSON THOMAS.
 XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX XX WPI; 2000-679325/66.
 XX DR
 XX PT New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer
 XX PS Claim 18; Page 17; 74pp; English.
 XX CC The invention relates to a peptide conjugate having the formula:
 CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the Bcl-2 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
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 Best Local Similarity 100.0%; Pred. No. 4.6e-14;
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 DB 1 NLWAAQRYGRELRLMSDEFGSFKGL 26
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 XX AC AAB37002;
 XX DT 28-FEB-2001 (first entry)
 XX DE Bcl2 polypeptide BH3 domain peptide #2.
 XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.
 XX OS Homo sapiens.
 XX WO200059526-A1.
 XX PN 12-OCT-2000.
 XX PD

PF 06-APR-2000; 2000WO-US09352.
 XX 07-APR-1999; 99US-0128202.
 XX PA (UYJE-) UNIV JEFFERSON THOMAS.
 XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX XX WPI; 2000-679325/66.
 XX DR
 XX PT New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer
 XX PS Claim 18; Page 17; 74pp; English.
 XX CC The invention relates to a peptide conjugate having the formula:
 CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
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 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the Bcl-2 domain of the cell death agonist Bad. The peptide conjugate is
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 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
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 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
 XX SQ Sequence 26 AA;
 Query Match 93.2%; Score 138; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 4.6e-14;
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 DB 1 NLWAAQRYGRELRLMSDEFGSFKGL 26
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 XX AC AAB37003;
 XX DT 28-FEB-2001 (first entry)
 XX DE Bcl2 polypeptide BH3 domain peptide #3.
 XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.
 XX OS Homo sapiens.
 XX WO200059526-A1.
 XX PN

XX 12-OCT-2000.
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 XX 06-APR-2000; 2000WO-US09352.
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 XX 07-APR-1999; 99US-0128202.
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 XX (UJVE-) UNIV JEFFERSON THOMAS.
 PA
 XX
 XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 PI
 XX
 XX WPI; 2000-679325/66.
 DR
 XX
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 PT
 XX
 XX Claim 18; Page 17; 7app; English.
 PS
 XX
 XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylene containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
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 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
 XX
 XX Sequence 27 AA;
 SQ
 Query Match 93.2%; Score 138; DB 21; Length 27;
 Best Local Similarity 100.0%; Pred. No. 4.8e-14;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 2 NLWAAQRYGRELRLMSDFEGSFKGL 27
 DB 1 NLWAAQRYGRELRLMSDFEGSFKGL 26
 RESULT 6
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 ID AAB70370 standard; protein; 162 AA. 1
 XX
 AC AAB70370;
 XX
 XX 02-MAY-2001 (first entry)
 DT
 XX Shorter murine BAD mutant amino acid sequence SEQ ID NO:3.
 DE
 XX Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 KW immunostimulant; neuroprotective; neurotropic; antiischemic; vulnary;
 KW cytostatic; antiviral; antiarthritic; antiinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.

XX Mus musculus.
 OS Synthetic.
 OS
 XX WO200110888-A1.
 PN
 XX 15-FEB-2001.
 PD
 XX 30-MAY-2000; 2000WO-US11864.
 PF
 XX 28-MAY-1999; 99US-0136783.
 PR
 XX (APOC-) APOPTOSIS TECHNOLOGY INC.
 PA
 XX Zhou X;
 PI
 XX WPI; 2001-138734/14.
 DR
 XX
 XX New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -
 PT
 XX Claim 7; Page 148-149; 157pp; English.
 PS
 XX The present invention describes an isolated or synthetic polypeptide
 CC (I) comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
 CC neurotropic, antiischemic, vulnary, cytostatic, antiviral,
 CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed shorter murine BAD mutant amino acid sequence from the present
 CC invention.
 XX
 XX Sequence 162 AA;
 SQ
 Query Match 93.2%; Score 138; DB 22; Length 162;
 Best Local Similarity 100.0%; Pred. No. 3.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDFEGSFKGL 27
 DB 98 NLWAAQRYGRELRLMSDFEGSFKGL 123
 RESULT 7
 AAR95168
 ID AAR95168 standard; Protein; 204 AA.
 XX
 AC AAR95168;
 XX
 XX 06-JAN-1997 (first entry)
 DT
 XX bcl-x(L)/bcl-2 associated death promoter protein.
 DE
 XX Epitope; murine; bcl-x(L)/bcl-2 associated death promoter; Bad; stroke;
 KW polypeptide; bcl-x; cell death; regulate; BH1; BH2; apoptotic cell death;
 KW cytokine deprivation; IL-3 dependent cell line; immunodeficiency; AIDS;
 KW neurodegenerative disease; senescence; ischaemia; neoplasia.
 XX
 XX Mus musculus.
 OS
 XX

```

FH Key Location/Qualifiers
FT Region 147..149
FT /note= "BH1 conserved amino acids"
FT Region 191..192
FT /note= "BH2 conserved amino acids"
FT Domain 38..61
FT /note= "PEST sequence"
FT Domain 111..130
FT /note= "PEST sequence"
XX WO9613614-A1.
XX
XX 09-MAY-1996.
XX
XX 31-OCT-1995; 95WO-US14246.
XX
XX 31-OCT-1994; 94US-0333565.
XX
XX (UNIW ) UNIV WASHINGTON.
XX
XX Korsmeyer SJ;
XX
XX WPI; 1996-251465/25.
XX N-PSDB; AAT29479.
XX
XX Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
XX useful to treat neoplasia and apoptosis and to identify agents
XX inhibiting its binding to bcl-2 or bcl-x(L) to form heteromultimers
XX
XX Claim 3; Fig 1; 130pp; English.
XX
XX This sequence represents the murine bcl-x(L)/bcl-2 associated death
XX promoter (Bad) gene. Bad is a 22.1 kD protein which interacts with
XX bcl-2 and bcl-x proteins and regulates cell death. It has homology
XX to the bcl-2-related family clustered in the BH1 and BH2 domain. Bad
XX has been found to hybridise to bcl-x(L) and bcl-2 in yeast two-hybrid
XX assays and in vivo in mammalian cells. Overexpressed Bad counters the
XX death inhibitory activity of bcl-x(L), but is much less effective at
XX countering the death inhibitory activity of bcl-2. Bad expression can
XX accelerate apoptotic cell death induced by cytokine deprivation in an
XX IL-3 dependent cell line expressing bcl-x(L), and it's also counters the
XX death repressor activity of bcl-x(L). Bad competes with Bax for binding
XX to bcl-x(L). Bad may be used to identify agents which inhibit its
XX binding to bcl-2 or bcl-x(L) to form heterodimers. Such agents may be
XX used to treat neurodegenerative diseases, immunodeficiency diseases,
XX e.g. AIDS, senescence or Ischaemia.
XX
XX Sequence 204 AA;
XX
XX Query Match 93.2%; Score 138; DB 17; Length 204;
XX Best Local Similarity 100.0%; Pred. No. 4.5e-13;
XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 2 NLWAAQRYGRELRRMSDEFGSPKGL 27
XX |
XX Db 140 NLWAAQRYGRELRRMSDEFGSPKGL 165
XX
XX RESULT 8
XX AAW61315
XX ID AAW61315 standard; Protein; 204 AA.
XX
XX AC AAW61315;
XX
XX DT 07-OCT-1998 (first entry)
XX
XX Murine BCL-XL/BCL-2 associated cell death regulator.
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
XX serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
XX
XX PN WO9817682-A1.

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PN WO9817682-A1.
XX
XX 30-APR-1998.
XX
XX 17-OCT-1997; 97WO-US19175.
XX
XX 18-OCT-1996; 96US-0733505.
XX
XX (UNIW ) UNIV WASHINGTON.
XX
XX Korsmeyer SJ;
XX
XX WPI; 1998-261422/23.
XX N-PSDB; AAV27833.
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
XX useful for, e.g. treating reduced apoptosis such as in cancer or
XX viral infection
XX
XX Claim 1; Fig 10; 95pp; English.
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
XX death regulator) proteins, having an amino acid other than Ser at
XX position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
XX present sequence is the murine BAD protein. Also described are: (1)
XX fragments of mutant BAD protein able to decrease cell viability; (2)
XX fusion proteins of mutant BAD with a heterologous polypeptide that
XX increases intracellular delivery. Mutant BAD proteins are used to treat
XX or prevent diseases associated with reduced apoptosis, e.g. cancer,
XX viral infection, lymphoproliferation, arthritis, infertility,
XX inflammation and autoimmune disease. Polynucleotide sequences encoding
XX mutant BAD proteins can be used similarly by gene therapy or to produce
XX transgenic animals for use as disease models or in drug screening. BAD
XX proteins phosphorylated at specified Ser are used to screen for enhancers
XX and inhibitors of serine-phosphatase. Inhibitors are potentially useful
XX in treatment of excessive apoptosis such as AIDS, neurodegeneration,
XX aging or ischaemic cell death. The apoptotic status of cells is
XX determined by measuring relative amounts of phosphorylated and non-
XX phosphorylated BAD, by usual immuncassays. Mutant BAD proteins have
XX greater death-promoting activity than wild-type BAD which can become
XX phosphorylated on the specified Ser, forming a product that does not
XX heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
XX proteins in the cytosol, thus promoting cell survival. The mutants with
XX Ser substituted cannot bind 14-3-3.
XX
XX Sequence 204 AA;
XX
XX Query Match 93.2%; Score 138; DB 19; Length 204;
XX Best Local Similarity 100.0%; Pred. No. 4.5e-13;
XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 2 NLWAAQRYGRELRRMSDEFGSPKGL 27
XX |
XX Db 140 NLWAAQRYGRELRRMSDEFGSPKGL 165
XX
XX RESULT 9
XX AAW61316
XX ID AAW61316 standard; Protein; 204 AA.
XX
XX AC AAW61316;
XX
XX DT 07-OCT-1998 (first entry)
XX
XX Mutant BCL-XL/BCL-2 associated cell death regulator #1.
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
XX serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
XX Synthetic.
XX
XX PN WO9817682-A1.

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XX 30-APR-1998.
PD
XX
XX 17-OCT-1997; 97WO-US19175.
PF
XX
XX 18-OCT-1996; 96US-0733505.
PR
XX
XX (UNIW ) UNIV WASHINGTON.
PA
XX
XX Korsmeyer SJ;
PI
XX
XX WPI; 1998-261422/23.
DR
XX
XX N-PSDB; AAV27834.
DR
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
PT useful for, e.g. treating reduced apoptosis such as in cancer or
PT viral infection
XX
XX
XX Claim 7; Page 59; 95pp; English.
PS
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
CC death regulator) proteins, having an amino acid other than Ser at
CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
CC present sequence represents a mutant BAD protein. Also described are: (1)
CC fragments of mutant BAD protein able to decrease cell viability; (2)
CC fusion proteins of mutant BAD with a heterologous polypeptide that
CC increases intracellular delivery. Mutant BAD proteins are used to treat
CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
CC viral infection, lymphoproliferation, arthritis, infertility,
CC inflammation and autoimmune disease. Polynucleotide sequences encoding
CC mutant BAD proteins can be used similarly by gene therapy or to produce
CC transgenic animals for use as disease models or in drug screening. BAD
CC proteins phosphorylated at specified Ser are used to screen for enhancers
CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
CC aging or ischaemic cell death. The apoptotic status of cells is
CC determined by measuring relative amounts of phosphorylated and non-
CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
CC greater death-promoting activity than wild-type BAD which can become
CC phosphorylated on the specified Ser, forming a product that does not
CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
CC proteins in the cytosol, thus promoting cell survival. The mutants with
CC Ser substituted cannot bind 14-3-3.
XX
XX
XX Sequence 204 AA;
SQ
Query Match 93.28; Score 138; DB 19; Length 204;
Best Local Similarity 100.0%; Pred. No. 4.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRELRRMSDEFFGSFKGL 27
DB 140 NLWAAQRYGRELRRMSDEFFGSFKGL 165
RESULT 10
AAW61317
ID AAW61317 standard; Protein; 204 AA.
XX
XX AAW61317;
AC
XX
XX 07-OCT-1998 (first entry)
DT
XX
XX Mutant BCL-XL/BCL-2 associated cell death regulator #2.
DE
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
KW serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
OS
XX Synthetic.
OS
XX W09817682-A1.
PN
XX
XX
XX

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PD 30-APR-1998.
XX
XX 17-OCT-1997; 97WO-US19175.
PF
XX
XX 18-OCT-1996; 96US-0733505.
PR
XX
XX (UNIW ) UNIV WASHINGTON.
PA
XX
XX Korsmeyer SJ;
PI
XX
XX WPI; 1998-261422/23.
DR
XX
XX N-PSDB; AAV27835.
DR
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
PT useful for, e.g. treating reduced apoptosis such as in cancer or
PT viral infection
XX
XX
XX Claim 7; Page 60; 95pp; English.
PS
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
CC death regulator) proteins, having an amino acid other than Ser at
CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
CC present sequence represents a mutant BAD protein. Also described are: (1)
CC fragments of mutant BAD protein able to decrease cell viability; (2)
CC fusion proteins of mutant BAD with a heterologous polypeptide that
CC increases intracellular delivery. Mutant BAD proteins are used to treat
CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
CC viral infection, lymphoproliferation, arthritis, infertility,
CC inflammation and autoimmune disease. Polynucleotide sequences encoding
CC mutant BAD proteins can be used similarly by gene therapy or to produce
CC transgenic animals for use as disease models or in drug screening. BAD
CC proteins phosphorylated at specified Ser are used to screen for enhancers
CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
CC aging or ischaemic cell death. The apoptotic status of cells is
CC determined by measuring relative amounts of phosphorylated and non-
CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
CC greater death-promoting activity than wild-type BAD which can become
CC phosphorylated on the specified Ser, forming a product that does not
CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
CC proteins in the cytosol, thus promoting cell survival. The mutants with
CC Ser substituted cannot bind 14-3-3.
XX
XX
XX Sequence 204 AA;
SQ
Query Match 93.28; Score 138; DB 19; Length 204;
Best Local Similarity 100.0%; Pred. No. 4.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRELRRMSDEFFGSFKGL 27
DB 140 NLWAAQRYGRELRRMSDEFFGSFKGL 165
RESULT 11
AAW61318
ID AAW61318 standard; Protein; 204 AA.
XX
XX AAW61318;
AC
XX
XX 07-OCT-1998 (first entry)
DT
XX
XX Mutant BCL-XL/BCL-2 associated cell death regulator #3.
DE
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
KW serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
OS
XX Synthetic.
OS
XX W09817682-A1.
PN
XX
XX
XX
XX 30-APR-1998.
PD

```


XX 17-OCT-1997; 97WO-US19175.
 XX 18-OCT-1996; 96US-0733505.
 XX (UNIW) UNIV WASHINGTON.
 XX Korsmeyer SJ;
 XX WPI; 1998-261422/23.
 DR N-PSDB; AAV27836.
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 XX Claim 7; Page 60-61; 95pp; English.
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX Sequence 204 AA;
 SQ Query Match 93.2%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
 DB 140 NLWAAQRYGRELRLMSDEFGSKGL 165
 RESULT 12
 AAW58832
 ID AAW58832 standard; protein; 204 AA.
 XX AAW58832;
 AC AAW58832;
 XX 23-JUL-1998 (first entry)
 DT Murine BAD protein.
 DE BAD protein; Bcl-XL/Bcl-2 associated cell death regulator; 14-3-3;
 KW serine phosphorylation; post-translational modification; apoptosis;
 KW signal transduction regulator; phosphoserine phosphatase; senescence;
 KW immunodeficiency disease; neurodegenerative disease; infertility;
 KW cancer, viral infection; lymphoproliferative condition; arthritis;
 KW inflammation; autoimmune diseases.
 XX Mus sp.
 OS Mus sp.
 XX WO9809643-A1.

XX 12-MAR-1998.
 PD 09-SEP-1997; 97WO-US15871.
 PF 09-SEP-1996; 96US-0707868.
 XX (UNIW) UNIV WASHINGTON.
 PA Korsmeyer SJ;
 XX WPI; 1998-207049/18.
 PI Serine-phosphorylated Bcl-XL/Bcl-2 Associated cell Death regulator
 DR polypeptide - useful for modulation of apoptosis associated with,
 XX e.g. cancer and immunodeficiency diseases
 XX Claim 3; Fig 8; 61pp; English.
 XX This sequence represents a novel serine-phosphorylated protein, BAD
 CC (Bcl-XL/Bcl-2 associated cell death regulator). The serine residue is
 CC phosphorylated in a post-translational modification and allows binding
 CC to the 14-3-3 protein which is a signal transduction regulator.
 CC Modulators of phosphorylated BAD, which act through inhibition/activation
 CC of a phosphoserine phosphatase, are useful for preventing/treating
 CC increased/decreased apoptosis in a cell. The increased apoptosis may
 CC result from immunodeficiency diseases, senescence, neurodegenerative and
 CC disease, ischaemic cell death, reperfusion cell death, infertility and
 CC wound-healing. Decreased apoptosis may result from cancer, viral
 CC infection, lymphoproliferative conditions, arthritis, infertility,
 CC inflammation and autoimmune diseases. Measuring the amount of
 CC phosphorylated compared to unphosphorylated BAD polypeptide and/or total
 CC BAD in a cell is useful for determining the apoptotic state of a cell.
 XX Sequence 204 AA;
 SQ Query Match 93.2%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
 DB 140 NLWAAQRYGRELRLMSDEFGSKGL 165
 RESULT 13
 AAB70369
 ID AAB70369 standard; protein; 204 AA.
 XX AAB70369;
 AC AAB70369;
 XX 02-MAY-2001 (first entry)
 DT Longer murine BAD mutant amino acid sequence SEQ ID NO:2.
 DE Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 XX immunostimulant; neuroprotective; nootropic; antischismatic; vulnary;
 KW cytostatic; antiviral; antiarthritic; antiinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.
 XX Mus musculus.
 OS Synthetic.
 OS WO200110888-A1.
 EN 15-FEB-2001.
 XX 30-MAY-2000; 2000WO-US11864.
 XX 28-MAY-1999; 99US-0136783.
 PR

XX (AFOP-) APOPTOSIS TECHNOLOGY INC.
 PA Zhou X;
 PI WPI; 2001-138734/14.
 DR
 XX New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -
 XX
 PS Claim 4; Page 148; 157pp; English.
 XX The present invention describes an isolated or synthetic polypeptide
 CC (1) comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (1) has immunostimulant, neuroprotective,
 CC neurotropic, antischaeamic, vulnerary, cytoskeletal, antiviral,
 CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed longer murine BAD mutant amino acid sequence from the present
 CC invention.
 XX
 SQ Sequence 204 AA;
 Query Match 93.2%; Score 138; DB 22; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSGKGL 27
 Db 140 NLWAAQRYGRELRLMSDEFGSGKGL 165
 RESULT 14
 ABR39082
 ID ABR39082 standard; Protein; 204 AA.
 XX ABR39082;
 AC
 XX 10-MAY-2003 (first entry)
 DT
 XX Murine BAD protein SEQ ID NO:4.
 DE
 XX Murine; BAD; herpes simplex virus; HSV; US3; herpes virus; apoptosis;
 KW virucide; infection.
 KW
 XX Mus musculus.
 OS
 XX WO2003012049-A2.
 PN
 XX 13-FEB-2003.
 PD
 XX 31-JUL-2002; 2002WO-US24177.
 PF
 XX 31-JUL-2001; 2001US-308929P.
 PR
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Munger J, Roizman B;
 PI WPI; 2003-248168/24.
 DR

DR N-PSDB; ABZ81201.
 XX Inducing apoptosis in a cell infected with herpes simplex virus, HSV,
 PT by administering to the cell, a composition comprising an agent that
 PT inhibits phosphorylation of pro-apoptotic polypeptide BAD by HSV US3 -
 XX
 PS Claim 15; Page 168; 192pp; English.
 XX The present invention describes a method (M1) for inducing apoptosis in
 CC a cell infected with herpes simplex virus (HSV), which comprises
 CC administering to the cell, a composition having an agent that inhibits
 CC phosphorylation of pro-apoptotic polypeptide BAD by HSV US3. Also
 CC described is a method (M2) for treating a patient infected with HSV, by
 CC administering to the patient, a composition comprising a peptide
 CC comprising a sequence of 4-100 continuous amino acids of a 168 residue
 CC amino acid sequence (see ABR39081), where the peptide comprises ser112,
 CC ser135, or ser155, or their combinations. BAD has virucide activity.
 CC M1 is useful for inducing apoptosis in a cell infected with HSV, where
 CC the cell is in a human. M2 is useful for treating a patient infected
 CC with HSV. The present sequence represents murine BAD, which is used in
 CC the exemplification of the present invention.
 XX
 SQ Sequence 204 AA;
 Query Match 93.2%; Score 138; DB 24; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSGKGL 27
 Db 140 NLWAAQRYGRELRLMSDEFGSGKGL 165
 RESULT 15
 AAU00220
 ID AAU00220 standard; Protein; 567 AA.
 XX AAU00220;
 AC
 XX 31-MAY-2001 (first entry)
 DT
 XX Bad-DTTR apoptosis-modifying fusion protein.
 DE
 XX Mouse; Bad-DTTR; apoptosis; cancer; spinal muscular atrophy;
 KW diphtheria toxin receptor binding domain; DTR; neoplasm; tumour;
 KW hyper-proliferation; Alzheimer's disease; neurodegenerative disorder;
 KW transient ischaemic neuronal injury; stroke; spinal cord injury;
 KW Huntington's disease.
 KW
 XX Chimeric - Mus sp.
 OS Chimeric - Corynebacterium diphtheriae.
 OS Chimeric - Synthetic.
 XX
 XX Key Location/Qualifiers
 FT Region 3..12
 FT /note= "10x histidine tag"
 XX
 PN WO200112661-A2.
 XX
 XX 22-FEB-2001.
 PD
 XX 15-AUG-2000; 2000WO-US22293.
 PF
 XX 16-AUG-1999; 99US-0149220.
 PR
 XX (HARD) HARVARD COLLEGE.
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 XX Youle RJ, Liu X, Collier RJ;
 PI WPI; 2001-218343/22.
 DR N-PSDB; AAS00248.
 XX

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:22:21 : Search time 14.6 Seconds
(without alignments)
81.144 Million cell updates/sec

Title: US-09-544-664-55

Perfect score: 148

Sequence: 1 KNLWAAQRYGRLRMSDFEGSPKGLX 28

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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1: /cgn2_6/ptodata/1/iaa/5A_COMB.pep.*
2: /cgn2_6/ptodata/1/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/1/iaa/8A_COMB.pep.*
4: /cgn2_6/ptodata/1/iaa/8B_COMB.pep.*
5: /cgn2_6/ptodata/1/iaa/PTUS_COMB.pep.*
6: /cgn2_6/ptodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	138	93.2	204	2	US-08-661-479-2
3	138	93.2	204	2	US-08-733-505A-1
4	138	93.2	204	2	US-08-733-505A-12
5	138	93.2	204	2	US-08-733-505A-13
6	138	93.2	204	2	US-08-733-505A-14
7	135	91.2	204	2	US-08-717-123-3
8	135	91.2	204	4	US-09-375-257-3
9	114	77.0	166	1	US-08-665-617-2
10	114	77.0	168	2	US-08-717-123-2
11	114	77.0	168	3	US-08-985-335-1
12	114	77.0	168	3	US-08-985-335-7
13	114	77.0	168	3	US-09-410-372-1
14	114	77.0	168	3	US-09-410-372-7
15	114	77.0	168	4	US-09-375-257-2
16	113	76.4	23	1	US-08-333-565-10
17	113	76.4	23	2	US-08-661-479-10
18	102	68.9	59	2	US-08-733-505A-55
19	102	68.9	59	2	US-08-733-505A-56
20	102	68.9	59	2	US-08-733-505A-57
21	102	68.9	59	2	US-08-733-505A-58
22	86	58.1	16	1	US-08-333-565-26
23	86	58.1	16	2	US-08-661-479-26
24	61	41.2	11	2	US-08-733-505A-34
25	61	41.2	11	2	US-08-706-741B-59
26	61	41.2	11	2	US-08-924-695A-69
27	51	34.5	66	2	US-08-867-087B-40

28 48.5 32.8 904 4 US-09-328-352-4656 Sequence 4656, Ap
29 46 31.1 610 4 US-09-252-991A-19594 Sequence 19594, A
30 46 31.1 946 3 US-09-074-579-3 Sequence 3, Appli
31 46 31.1 946 3 US-09-388-774-3 Sequence 3, Appli
32 46 31.1 946 4 US-09-546-153-1 Sequence 1, Appli
33 45.5 30.7 906 4 US-09-252-991A-31458 Sequence 31458, A
34 45 30.4 229 4 US-09-252-991A-23807 Sequence 23807, A
35 45 30.4 303 4 US-09-328-352-5164 Sequence 5164, Ap
36 45 30.4 356 4 US-09-235-103-2 Sequence 2, Appli
37 45 30.4 356 4 US-09-235-103-4 Sequence 4, Appli
38 45 30.4 1064 4 US-09-252-991A-17508 Sequence 17508, A
39 44.5 30.1 903 4 US-09-252-991A-28775 Sequence 28775, A
40 44 29.7 125 4 US-09-328-352-7449 Sequence 7449, Ap
41 44 29.7 263 4 US-09-651-656-27 Sequence 27, Appl
42 44 29.7 263 4 US-09-650-855-27 Sequence 27, Appl
43 44 29.7 277 4 US-09-252-991A-28581 Sequence 28581, A
44 44 29.7 877 4 US-09-206-551-20 Sequence 20, Appl
45 44 29.7 1125 4 US-09-252-991A-18729 Sequence 18729, A

ALIGNMENTS

RESULT 1
US-08-333-565-2
; Sequence 2, Application US/083333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/333,565
; APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "Deduced amino acid sequence
; OTHER INFORMATION: of mouse BAD."
US-08-333-565-2

Query Match 93.2%; Score 138; DB 1; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSFKGL 27
140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 2
US-08-661-479-2
; Sequence 2, Application US/08661479
; Patent No. 5834209
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/661,479
; FILING DATE: 11-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "Deduced amino acid sequence
; OF mouse BAd."

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSFKGL 27
140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 3
US-08-733-505A-1
; Sequence 1, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TELECOMMUNICATION INFORMATION:
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:

; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-733-505A-1

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSFKGL 27
140 NLWAAQRYGRELRRMSDFEGSFKGL 165

RESULT 4
US-08-733-505A-12
; Sequence 12, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TELECOMMUNICATION INFORMATION:
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 12:

SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-12

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 5
US-08-733-505A-13
Sequence 13, Application US/08733505A
Patent No. 5856445
GENERAL INFORMATION:
APPLICANT: KORSMEYER, STANLEY J.
TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
NUMBER OF SEQUENCES: 60
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOWELL & HAFERKAMP, L.C.
STREET: 7733 FORSYTH BLVD., SUITE 1400
CITY: ST. LOUIS
STATE: MISSOURI
COUNTRY: USA
ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-13

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 6
US-08-733-505A-14
Sequence 14, Application US/08733505A
Patent No. 5856445
GENERAL INFORMATION:
APPLICANT: KORSMEYER, STANLEY J.

TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
NUMBER OF SEQUENCES: 60
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOWELL & HAFERKAMP, L.C.
STREET: 7733 FORSYTH BLVD., SUITE 1400
CITY: ST. LOUIS
STATE: MISSOURI
COUNTRY: USA
ZIP: 63105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/733,505A
FILING DATE:
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-14

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
Db 140 NLWAAQRYGRELRLMSDEFGSKGL 165

RESULT 7
US-08-717-123-3
Sequence 3, Application US/08717123
Patent No. 5965703
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
Acids and Methods of Use
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell and Flores
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: United States
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/717,123
FILING DATE: 20-SEP-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815

REFERENCE/DOCKET NUMBER: P-ID 1929
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
TOPOLOGY: linear
US-08-717-123-3

Query Match 91.2%; Score 135; DB 2; Length 204;
Best Local Similarity 96.2%; Pred. No. 4.2e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
|||||
Db 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 8

US-09-375-257-3
Sequence 3, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:

APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
FILE REFERENCE: 480140.428D1
CURRENT APPLICATION NUMBER: US/09/375,257
CURRENT FILING DATE: 1999-08-16
NUMBER OF SEQ ID NOS: 15
SOFTWARE: FastSeq for Windows version 4.0
SEQ ID NO 3
LENGTH: 204
TYPE: PRT
ORGANISM: Mus musculus
US-09-375-257-3

Query Match 91.2%; Score 135; DB 4; Length 204;
Best Local Similarity 96.2%; Pred. No. 4.2e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
|||||
Db 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 9

US-08-665-617-2
Sequence 2, Application US/08665617
Patent No. 5663316
GENERAL INFORMATION:

APPLICANT: Xu, Dong, Yin
TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Saliwanchik & Saliwanchik
STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: Florida
COUNTRY: USA
ZIP: 32606
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/665,617
FILING DATE:
CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:
NAME: Saliwanchik, David R.
REGISTRATION NUMBER: 31,794
REFERENCE/DOCKET NUMBER: CL-8
TELECOMMUNICATION INFORMATION:
TELEPHONE: (352) 375-8100
TELEFAX: (352) 372-5800
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 166 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-665-617-2

Query Match 77.0%; Score 114; DB 1; Length 166;
Best Local Similarity 91.7%; Pred. No. 5.4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKF 25
|||||
Db 101 NLWAAQRYGRELRRMSDEFVDSFK 124

RESULT 10

US-08-717-123-2
Sequence 2, Application US/08717123
Patent No. 5985703
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
TITLE OF INVENTION: Acids and Methods of Use
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell and Flores
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: United States
ZIP: 92122
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/717,123
FILING DATE: 20-SEP-1996
CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-ID 1929
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-717-123-2

Query Match 77.0%; Score 114; DB 2; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKF 25
|||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

```

CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/985,335
FILING DATE: Filed Herewith
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0421 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 1683637
US-08-985-335-7

Query Match 77.08; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred.No.5.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels

QY 2 NLWAAQRYGRELRRMSDEFGSPK 25
DB 103 NLWAAQRYGRELRRMSDEFGVDSFK 126
|||||
|||||

RESULT 13
US-09-410-372-1
Sequence 1, Application US/09410372
Patent No. 6281334
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
PROLIFERATION
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/410,372
FILING DATE:

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;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 08/985,335
;; FILING DATE:
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Billings, Lucy J.
;; REGISTRATION NUMBER: 36,749
;; REFERENCE/DOCKET NUMBER: PF-0421 US
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 650-855-0555
;; TELEFAX: 650-845-4166
;; INFORMATION FOR SEQ ID NO: 1:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 168 amino acids
;; TYPE: amino acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; IMMEDIATE SOURCE:
;; LIBRARY: SYNORAB01
;; CLONE: 358673
US-09-410-372-1

Query Match 77.0%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFEGSEK 25
|||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 14
US-09-410-372-7
;; Sequence 7, Application US/09410372
;; Patent No. 6281334
;; GENERAL INFORMATION:
;; APPLICANT: Hillman, Jennifer L.
;; APPLICANT: Yue, Henry
;; APPLICANT: Lal, Preeti
;; APPLICANT: Shah, Purvi
;; APPLICANT: Corley, Neil C.
;; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
;; TITLE OF INVENTION: PROLIFERATION
;; NUMBER OF SEQUENCES: 9
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Incyte Pharmaceuticals, Inc.
;; STREET: 3174 Porter Dr.
;; CITY: Palo Alto
;; STATE: CA
;; COUNTRY: USA
;; ZIP: 94304
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Diskette
;; COMPUTER: IBM Compatible
;; OPERATING SYSTEM: DOS
;; SOFTWARE: FastSeq for Windows Version 2.0
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/09/410,372
;; FILING DATE:
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 08/985,335
;; FILING DATE:
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Billings, Lucy J.
;; REGISTRATION NUMBER: 36,749
;; REFERENCE/DOCKET NUMBER: PF-0421 US
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 650-855-0555
;; TELEFAX: 650-845-4166
;; INFORMATION FOR SEQ ID NO: 7:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 168 amino acids
;; TYPE: amino acid
;; STRANDEDNESS: single

;; TOPOLOGY: linear
;; IMMEDIATE SOURCE:
;; LIBRARY: GenBank
;; CLONE: 1683637
US-09-410-372-7

Query Match 77.0%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFEGSEK 25
|||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 15
US-09-375-257-2
;; Sequence 2, Application US/09375257
;; Patent No. 6504022
;; GENERAL INFORMATION:
;; APPLICANT: Horne, William A.
;; APPLICANT: Oltersdorf, Tilman
;; TITLE OF INVENTION: HUMAN RAD POLYPEPTIDES, ENCODING NUCLEIC
;; TITLE OF INVENTION: ACIDS AND METHODS OF USE
;; FILE REFERENCE: 480140.428D1
;; CURRENT APPLICATION NUMBER: US/09/375,257
;; CURRENT FILING DATE: 1999-08-16
;; NUMBER OF SEQ ID NOS: 15
;; SOFTWARE: FastSeq for Windows Version 4.0
;; SEQ ID NO 2
;; LENGTH: 168
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-09-375-257-2

Query Match 77.0%; Score 114; DB 4; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.5e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFEGSEK 25
|||||
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

Search completed: September 15, 2003, 17:45:06
Job time : 14.6 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:25:56 ; Search time 22.2 Seconds
(without alignments)
184.034 Million cell updates/sec

Title: US-09-544-664-55
Perfect score: 148
Sequence: 1 NLWAAQRYGRLRRMSDEFGSKGLK 28

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 541936 seqs, 145912426 residues
Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA:*

1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
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11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
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14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	135	91.2	204	9 US-09-922-378-3	Sequence 3, Appli
2	135	91.2	204	14 US-10-066-179-3	Sequence 3, Appli
3	114	77.0	25	15 US-10-059-261-258	Sequence 258, App
4	114	77.0	168	9 US-09-922-378-2	Sequence 2, Appli
5	114	77.0	168	9 US-09-894-657-1	Sequence 1, Appli
6	114	77.0	168	9 US-09-894-657-7	Sequence 7, Appli
7	114	77.0	168	14 US-10-066-179-2	Sequence 2, Appli
8	71	48.0	15	15 US-10-174-1058-147	Sequence 147 App
9	52	35.1	215	15 US-10-156-761-9145	Sequence 9145, Ap
10	51	34.5	682	12 US-10-238-075-1077	Sequence 1077, Ap
11	47	31.8	35	15 US-10-092-750-1	Sequence 1, Appli
12	47	31.8	138	15 US-10-092-750-241	Sequence 241, App
13	46	31.1	946	9 US-09-828-423-3	Sequence 3, Appli
14	44	29.7	162	11 US-09-934-455-162	Sequence 162, App
15	44	29.7	272	15 US-10-156-761-11541	Sequence 11541, A

16	44	29.7	426	9 US-09-815-242-5704	Sequence 5704, Ap
17	44	29.7	699	14 US-10-008-355-8	Sequence 8, Appli
18	44	29.7	705	9 US-09-815-242-12463	Sequence 12463, A
19	44	29.7	712	14 US-10-008-355-2	Sequence 2, Appli
20	44	29.7	877	12 US-10-369-294-20	Sequence 20, Appli
21	43	29.1	94	9 US-09-864-761-39540	Sequence 39540, A
22	43	29.1	164	11 US-09-986-480-395	Sequence 395, App
23	43	29.1	213	9 US-09-843-846-18	Sequence 18, Appli
24	43	29.1	232	10 US-09-881-752A-238	Sequence 238, App
25	43	29.1	380	9 US-09-149-045-2	Sequence 2, Appli
26	43	29.1	380	15 US-10-166-359-2	Sequence 2, Appli
27	43	29.1	380	15 US-10-166-113-2	Sequence 2, Appli
28	43	29.1	380	15 US-10-166-357-2	Sequence 2, Appli
29	43	29.1	380	15 US-10-166-372-2	Sequence 2, Appli
30	43	29.1	380	15 US-10-184-722-3	Sequence 3, Appli
31	43	29.1	380	15 US-10-251-385-62	Sequence 62, Appli
32	43	29.1	380	15 US-10-251-385-198	Sequence 198, App
33	43	29.1	380	15 US-10-225-567A-233	Sequence 233, App
34	43	29.1	543	15 US-10-156-761-13485	Sequence 13485, A
35	43	29.1	571	9 US-09-815-242-11813	Sequence 11813, A
36	43	29.1	582	10 US-09-331-631A-22	Sequence 22, Appli
37	43	29.1	640	9 US-09-989-722-501	Sequence 501, App
38	43	29.1	640	9 US-09-989-723-501	Sequence 501, App
39	43	29.1	640	9 US-09-989-279-501	Sequence 501, App
40	43	29.1	640	9 US-09-989-727-501	Sequence 501, App
41	43	29.1	640	10 US-09-989-731-501	Sequence 501, App
42	43	29.1	640	10 US-09-989-732-501	Sequence 501, App
43	43	29.1	640	10 US-09-991-073-501	Sequence 501, App
44	43	29.1	640	10 US-09-909-320-292	Sequence 292, App
45	43	29.1	640	10 US-09-990-442-501	Sequence 501, App

ALIGNMENTS

RESULT 1
US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 91.2%; Score 135; DB 9; Length 204;
Best local Similarity 96.2%; Pred. No. 8.2e-12;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 2 NLWAAQRYGRLRRMSDEFGSKGL 27
Db 140 NLWAAQRYGRLRRMTDFEFGSKGL 165

RESULT 2
US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE

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; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066.179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
; US-10-066-179-3

Query Match          91.2%; Score 135; DB 14; Length 204;
Best Local Similarity 96.2%; Pred. No. 8.2e-12;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
   |||||
Db 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 3
US-10-059-261-258
; Sequence 258, Application US/10059261
; Publication No. US20030077826A1
; GENERAL INFORMATION:
; APPLICANT: EDELMAN, LENA
; APPLICANT: JACOTOT, ETIENNE DANIEL FRANCOIS
; APPLICANT: BRAND, JEAN-PAUL
; TITLE OF INVENTION: CHIMERIC MOLECULES CONTAINING A MODULE ABLE TO TARGET
; TITLE OF INVENTION: SPECIFIC CELLS AND A MODULE REGULATING THE APOPTOTIC
; TITLE OF INVENTION: FUNCTION OF THE PERMEABILITY TRANSITION PORE COMPLEX
; TITLE OF INVENTION: (PTPC)
; FILE REFERENCE: 03495.0216
; CURRENT APPLICATION NUMBER: US/10/059,261
; CURRENT FILING DATE: 2002-08-29
; PRIOR APPLICATION NUMBER: 60/265,594
; PRIOR FILING DATE: 2001-02-02
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: Patent In Ver. 2.1
; SEQ ID NO 258
; LENGTH: 25
; TYPE: PRT
; ORGANISM: Unknown Organism
; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: TOX peptide
; US-10-059-261-258

Query Match          77.0%; Score 114; DB 15; Length 25;
Best Local Similarity 91.7%; Pred. No. 1e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKF 25
   |||||
Db 1 NLWAAQRYGRELRRMSDEFGSKF 24

RESULT 4
US-09-922-378-2
; Sequence 2, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
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; ORGANISM: Homo sapiens
; US-09-922-378-2

Query Match          77.0%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 7.2e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKF 25
   |||||
Db 103 NLWAAQRYGRELRRMSDEFGSKF 126

RESULT 5
US-09-894-657-1
; Sequence 1, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
; US-09-894-657-1

Query Match          77.0%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 7.2e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKF 25
   |||||
Db 103 NLWAAQRYGRELRRMSDEFGSKF 126

RESULT 6
US-09-894-657-7
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; Sequence 7, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Inventors: Yue, Henry
;           Lal, Preeti
;           Shah, Purvi
;           Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
;                 PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 7:
US-09-894-657-7

Query Match          77.0%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 7.2e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2 NLWAAQRYGRELRRMSDEFGSFK 25
Db      103 NLWAAQRYGRELRRMSDEFDVDFK 126

RESULT 7
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168

; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-066-179-2

Query Match          77.0%; Score 114; DB 14; Length 168;
Best Local Similarity 91.7%; Pred. No. 7.2e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2 NLWAAQRYGRELRRMSDEFGSFK 25
Db      103 NLWAAQRYGRELRRMSDEFDVDFK 126

RESULT 8
US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US20030068652A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECIF
; IC ANTIBODIES COUPLED WITH DATABASE SEARCHIN
; FILE REFERENCE: CST-138 CIP3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147

Query Match          48.0%; Score 71; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.00097;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      10 GRELRMSDEFGS 23
Db      1 GRELRMSDEFGS 14

RESULT 9
US-10-156-761-9145
; Sequence 9145, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
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; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 9145
; LENGTH: 215
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-9145

Query Match          35.1%; Score 52; DB 15; Length 215;
Best Local Similarity 45.5%; Pred. No. 8.2;
Matches 10; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY      7 QRYGRLRMSDFEGSGFKLK 28
       :|:|||||||:|:|
Db      108 ERWGGDLRRMRDEADKYPELR 129

RESULT 10
US-10-238-075-1077
; Sequence 1077, Application US/10238075
; Publication No. US20030148324A1
; GENERAL INFORMATION:
; APPLICANT: I.N.S.E.R.M.
; TITLE OF INVENTION: Polynucleotides which are of nature B2/D- A- and which are isolated from
; FILE REFERENCE: BLANDINE
; CURRENT APPLICATION NUMBER: US/10/238,075
; CURRENT FILING DATE: 2002-09-10
; PRIOR APPLICATION NUMBER: 0003145
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 1576
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 1077
; LENGTH: 682
; TYPE: PRT
; ORGANISM: Escherichia coli
US-10-238-075-1077

Query Match          34.5%; Score 51; DB 12; Length 682;
Best Local Similarity 40.7%; Pred. No. 37;
Matches 11; Conservative 5; Mismatches 11; Indels 0; Gaps 0;

QY      1 KNLWAQRVGRLELRMSDFEGSGFKGL 27
       |:|||||:|:|:|
Db      608 KQTIAAQANGAKVPVRNRNGFTSMDIGL 634

RESULT 11
US-10-092-750-1
; Sequence 1, Application US/10092750
; Publication No. US20030032157A1
; GENERAL INFORMATION:
; APPLICANT: Hammond, Philip W.
; APPLICANT: Alpin, Julia
; TITLE OF INVENTION: Polypeptides Interactive with BCL-Xl
; FILE REFERENCE: 50036/050002
; CURRENT APPLICATION NUMBER: US/10/092,750
; CURRENT FILING DATE: 2002-03-07
; PRIOR APPLICATION NUMBER: US 60/274,526
; PRIOR FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 253
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 35
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-092-750-1

Query Match          31.8%; Score 47; DB 15; Length 35;
Best Local Similarity 45.5%; Pred. No. 6.7;
Matches 10; Conservative 3; Mismatches 5; Indels 4; Gaps 1;

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; TELEX: <Unknown>
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 946 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GENEBANK
; CLONE: g133985
; SEQUENCE DESCRIPTION: SEQ ID NO: 3 :
US-09-828-423-3

Query Match 31.1%; Score 46; DB 9; Length 946;
Best Local Similarity 30.8%; Pred. No. 2.7e+02;
Matches 8; Conservative 5; Mismatches 13; Indels 0; Gaps 0;

Qy 2 NLWAAQRYGRELRRMSDEFEGSKGL 27
Db 212 DWWIEPGLFELHVPDTFEGHEDGV 237

RESULT 14
US-09-934-455-162
; Sequence 162, Application US/09934455
; Publication No. US20030121070A1
; GENERAL INFORMATION:
; APPLICANT: Adam, Luc
; APPLICANT: Creelman, Robert
; APPLICANT: Dubell, Arnold
; APPLICANT: Heard, Jacqueline
; APPLICANT: Jiang, Cai-Zhong
; APPLICANT: Keddie, James
; APPLICANT: Pilgrim, Marsha
; APPLICANT: Ratcliffe, Oliver
; APPLICANT: Reuber, Lynne
; APPLICANT: Riechmann, Jose Luis
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Pineda, Onaira
; TITLE OF INVENTION: Genes for Modifying Plant Traits IV
; FILE REFERENCE: MBI-0025
; CURRENT APPLICATION NUMBER: US/09/934,455
; CURRENT FILING DATE: 2001-08-22
; PRIOR FILING DATE: 60/227439
; PRIOR FILING DATE: 2000-08-22
; PRIOR APPLICATION NUMBER: MBI-0022
; PRIOR FILING DATE: 2001-11-16
; PRIOR APPLICATION NUMBER: MBI-0023
; PRIOR FILING DATE: 2001-04-17
; NUMBER OF SEQ ID NOS: 516
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 162
; LENGTH: 270
; TYPE: PRT
; ORGANISM: Arabidopsis thaliana
US-09-934-455-162

Query Match 29.7%; Score 44; DB 11; Length 270;
Best Local Similarity 40.7%; Pred. No. 4.5e+02;
Matches 11; Conservative 4; Mismatches 8; Indels 4; Gaps 1;

Qy 2 NLWAAQRYGRELRRMSDEFEGSKGLK 28
Db 78 NMFQPIYGRDFKRSS----SSMVGLK 100

RESULT 15
US-10-156-761-11541
; Sequence 11541, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 11541
; LENGTH: 272
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-11541

Query Match 29.7%; Score 44; DB 15; Length 272;
Best Local Similarity 53.3%; Pred. No. 1.5e+02;
Matches 8; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

Qy 4 WAAQRYGELRRMSD 18
Db 29 WIAAHGAELELRAD 43

Search completed: September 15, 2003, 17:47:53
Job time : 22.2 secs
```

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:18:16 : Search time 12.6 Seconds
(without alignments)
213.708 Million cell updates/sec

Title: US-09-544-664-55
Perfect score: 148
Sequence: 1 KNLWAAQRYGRELRRMSDEFGSPKGLK 28

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues
Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_76:*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	138	93.2	204	A55671	bad protein - mous
2	54	36.5	946	JC5575	inter-alpha-trypsi
3	53	35.8	223	D70760	hypothetical prote
4	53	35.8	946	S54354	inter-alpha-inhibi
5	52	35.1	370	S38185	2-dehydro-3-deoxy-
6	51	34.5	232	A42095	floral homeotic pr
7	50	33.8	374	C84338	spermidine/putresc
8	50	33.8	516	A96753	probable threonine
9	50	33.8	1378	A81393	DNA-directed RNA p
10	49.5	33.4	127	A11210	glycerol-3-phospha
11	49	33.1	453	E83517	conserved hypotet
12	48.5	32.8	134	S40376	Ig kappa chain - h
13	48.5	32.8	514	T02975	annexin P35 - maiz
14	48	32.4	206	C36365	transforming prote
15	48	32.4	220	F72289	oxidoreductase, so
16	48	32.4	526	T08545	threonine synthase
17	48	32.4	1164	T24806	hypothetical prote
18	47.5	32.1	334	A39172	Antho-Ramide neur
19	47.5	32.1	1140	T09486	hypothetical prote
20	47	31.8	287	S43852	neuropeptide Pol-R
21	47	31.8	597	T82308	oxaloacetate decar
22	47	31.8	967	R26668	oxoglutarate dehyd
23	47	31.8	5138	T96895	hypothetical prote
24	46.5	31.4	514	T02961	annexin P33 - maiz
25	46.5	31.4	435	A44308	Antho-Ramide prec
26	46	31.1	165	S59899	chlorocruorin chai
27	46	31.1	399	T35440	probable polyamine
28	46	31.1	946	1YH02	inter-alpha-trypsi
29	45.5	30.7	261	G69510	conserved hypotet

30	45.5	30.7	327	2	AF2859	conserved hypotet
31	45.5	30.7	327	2	D97636	probable secreted
32	45.5	30.7	562	2	C71473	hypothetical prote
33	45.5	30.7	905	2	G83314	NADH dehydrogenase
34	45.5	30.7	1014	2	T36031	excinuclease ABC c
35	45	30.4	273	2	S06736	photosystem II oxy
36	45	30.4	273	2	AG2287	manganese-stabilizi
37	45	30.4	295	2	P83201	conserved hypotet
38	45	30.4	346	2	H93406	conserved hypotet
39	45	30.4	486	2	T31294	hypothetical prote
40	45	30.4	591	2	B44465	sodium ion pump ox
41	45	30.4	591	2	AB0509	oxaloacetate decar
42	45	30.4	591	2	AE0909	oxaloacetate decar
43	45	30.4	596	2	A28088	oxaloacetate decar
44	45	30.4	715	2	S52675	probable membrane
45	45	30.4	864	1	VCLJG4	env polyprotein -

ALIGNMENTS

RESULT 1

A55671
bad protein - mouse
C:Species: Mus musculus (house mouse)
C>Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999
C:Accession: A55671
Cell 80, 285-291, 1995
A>Title: Bad, a heterodimeric partner for Bcl-x-L and Bcl-2, displaces Bax and promot
A:Reference number: A55671; MUID:95136361; PMID:7834748
A:Accession: A55671
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-204 <YAN>
A:Cross-references: GB:L37296; NID:g639778; PIDN:AAA64465.1; PID:g639779
C:Keywords: heterodimer

Query Match 93.2%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.2e-12;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSPKGL 27
DB 140 NLWAAQRYGRELRRMSDEFGSPKGL 165
|||||

RESULT 2

JC5575
inter-alpha-trypsin inhibitor heavy chain 2 - golden hamster
C:Species: Mesocricetus auratus (golden hamster)
C>Date: 23-Sep-1997 #sequence_revision 23-Sep-1997 #text_change 20-Jun-2000
C:Accession: JC5575; PC4485
R:Nakatani, T.; Suzuki, Y.; Yamamoto, T.; Sinohara, H.
J. Biochem. 122, 71-82, 1997
A>Title: Molecular cloning and sequencing of cDNAs encoding three heavy-chain precurs
sin inhibitor heavy chain family.
A:Reference number: JC5574; MUID:97420688; PMID:9276673
A:Accession: JC5575
A:Molecule type: mRNA
A:Residues: 1-946 <NAX>
A:Cross-references: DDBJ:D89286; NID:g1694689; PIDN:BAAL3939.1; PID:g1694690
A:Experimental source: liver
A:Accession: PC4485
A:Molecule type: protein
A:Residues: 55-64;140-146;151-156;424-447;500-528;577-605 <NA2>
C:Comment: In the plasma three inter-alpha-trypsin inhibitor heavy chains 1, 2 and 3
that the complexes play important role for pancreatic cancer.
C:Superfamily: inter-alpha-trypsin inhibitor complex component II
F:261-264,717-916/Disulfide bonds: #status predicted
Query Match 36.5%; Score 54; DB 2; Length 946;
Best Local Similarity 34.6%; Pred. No. 11;

A;Residues: 1-232 <JAC>
A;Cross-references: GB:M86357; NID:g166607; PIDN:AAA32740.1; PID:g166608
A;Experimental source: petals, stamens
A;Note: sequence extracted from NCBI backbone (NCBIN:82520, NCBIIP:82521)
R;Okamoto, H.; Yano, A.; Shirasahi, H.; Okada, K.; Shimura, Y.
Plant Mol. Biol. 26, 465-472, 1994
A;Title: Genetic complementation of a floral homeotic mutation, apetala3, with an Arabidopsis gene
A;Reference number: S52633; MUID:95036018; PMID:7948893
A;Accession: S52633
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-63 <OKA>
A;Cross-references: GB:D21125
R;Blecker, H.; Mewes, H.W.; Lemcke, K.; Mayer, K.F.X.; Quetier, F.; Salanoubat M.Mewes, submitted to the Protein Sequence Database, March 2000
A;Reference number: Z24469
A;Accession: T47593
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-232 <BLO>
A;Cross-references: EMBL:AL132971
A;Experimental source: cultivar Columbia; BAC clone T12E18
C;Genetics:
A;Map position: 3
A;Introns: 63/2; 85/3; 106/2; 139/3; 153/3; 168/3
A;Note: T12E18.30
C;Superfamily: transcription factor squa; serum response factor DNA-binding domain homolog
C;Keywords: DNA binding; nucleus; transcription regulation
F:2-57/Domain: serum response factor DNA-binding domain homology <SRF>
Query Match 34.5%; Score 51; DB 2; Length 232;
Best Local Similarity 44.4%; Pred. No. 7.3;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;
QY 7 QRYG-----RELNRMSDEFEGSK 25
||| ||||| ||| |||
Db 107 QRLGCELDLDLQELRLDEMENTFK 133
RESULT 7
C84338
A;Title: spermidine/putrescine ABC transporter [imported] - Halobacterium sp. NRC-1
C;Species: Halobacterium sp. NRC-1
C;Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C;Accession: C84338
R;Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.; Lethausner, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jabloch, Jung, K.H.; Alam, M.; Freitas, T.
Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000
A;Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li
A;Title: Genome sequence of Halobacterium species NRC-1
A;Reference number: A84160; MUID:20504483; PMID:11016950
A;Accession: C84338
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-374 <STO>
A;Cross-references: GB:AE004437; NID:g10581314; PIDN:ARG20071.1; GSPDB:GN00138
C;Genetics:
A;Gene: potA2
Query Match 33.8%; Score 50; DB 2; Length 374;
Best Local Similarity 76.9%; Pred. No. 17;
Matches 10; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 12 ELRRMSDEFEGSF 24
||||| |||||
Db 197 ELRLSDAVEGSF 209
RESULT 8
A96753
A;Title: probable threonine synthase [imported] - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cross)
C;Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C;Accession: A96753
R;Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Aloni, Chiu, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Dewar, A.R.; Creasy, T.H.; Dewar, A.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A;Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim, C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marziani, Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A;Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon, ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A;Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A;Reference number: A86141; MUID:21016719; PMID:11130712
A;Accession: A96753
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-516 <STO>
A;Cross-references: GB:AE005173; NID:g5903070; PIDN:AAD55628.1; GSPDB:GN00141
C;Genetics:
A;Gene: F3N23.1
A;Map position: 1
Query Match 33.8%; Score 50; DB 2; Length 516;
Best Local Similarity 35.3%; Pred. No. 23;
Matches 12; Conservative 7; Mismatches 7; Indels 8; Gaps 1;
QY 2 NLWAAQRYGRELNRMSD-----EFEGSFKGL 27
||| ||||| ||| |||
Db 163 NLWAAERFGQYLMQNDLWVKHGISTGSKDL 196
RESULT 9
A81393
A;Title: DNA-directed RNA polymerase (EC 2.7.7.6) beta chain Cj0478 [imported] - Campylobacter.
C;Species: Campylobacter jejuni
C;Date: 31-Mar-2000 #sequence_revision 31-Mar-2000 #text_change 03-Jun-2002
C;Accession: A81393
R;Parkhill, J.; Wren, B.W.; Mungall, K.; Ketley, J.M.; Churcher, C.; Basham, D.; Chis, C.W.; Quail, M.; Rajandream, M.A.; Rutherford, K.M.; Vanvliet, A.; Whitehead, S.; Bann, Nature 403, 665-668, 2000
A;Title: The genome sequence of the food-borne pathogen Campylobacter jejuni reveals
A;Reference number: A81250; MUID:20150912; PMID:10689204
A;Accession: A81393
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-1378 <PAR>
A;Cross-references: GB:AL139075; GB:AL111168; NID:g6967817; PIDN:CA875116.1; PID:g696999
A;Experimental source: serotype O2, strain NCTC 11168
C;Genetics:
A;Gene: rpoB; Cj0478
C;Superfamily: DNA-directed RNA polymerase beta chain
C;Keywords: nucleotidyltransferase
Query Match 33.8%; Score 50; DB 2; Length 1378;
Best Local Similarity 40.6%; Pred. No. 63;
Matches 13; Conservative 3; Mismatches 10; Indels 6; Gaps 2;
QY 3 LWAAQRYG--RELNR-----SDEFECSFKGLX 28
||| ||| ||| |||
Db 1306 VWALEAYGAHTLREMLTIKSDVDEGRFSAYK 1337
RESULT 10
A11210
A;Title: glycerol-3-phosphate cytidyltransferase (gct), CDP-glycerol pyrophosphorylase (te1
C;Species: Listeria monocytogenes
C;Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 01-Mar-2002
C;Accession: A11210
R;Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloe, D.; Dominguez-Bernal, G.; Buchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, Science 294, 849-852, 2001
A;Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.

R; Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwinn, M.L.; Dodson, R.J.; Haft, D.H.; Hickey, J.A.; Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson, D.; C.M.

Nature 399, 323-329, 1999

A; Title: Evidence for lateral gene transfer between Archaea and Bacteria from genome sequence of *Mycobacterium thermophilus*

A; Reference number: A72200; PMID:9287316; PMID:10360571

A; Accession: F72289

A; Status: preliminary

A; Molecule type: DNA

A; Residues: 1-220 <ARN>

A; Cross-references: GB:AE001772; GB:AE000512; NID:94981693; PIDN:AD36230.1; PID:9498170

A; Experimental source: strain MSB8

C; Genetics:

A; Gene: TM1154

C; Superfamily: yeast SOL3 protein

Query Match 32.4%; Score 48; DB 2; Length 220;

Best Local Similarity 34.8%; Pred. NO. 19;

Matches 8; Conservative 8; Mismatches 7; Indels 0; Gaps 0;

QY 5 AAQRYGRELRLMSDEFESEFKGL 27

Db 111 ACEKYEIRSATDQFDLAILGM 133

Search completed: September 15, 2003, 17:27:03

Job time : 12.6 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:55 ; Search time 6.6 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-55

Perfect score: 148

Sequence: 1 KNLWAAQRYGRELRLMSDEPGSKGLK 28

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	138	93.2	204	1	BAD_MOUSE
2	138	93.2	205	1	BAD_RAT
3	114	77.0	168	1	BAD_HUMAN
4	54	36.5	946	1	ITH2_MESAU
5	53	35.8	946	1	ITH2_MOUSE
6	52.5	35.5	506	1	MATK_LEDPA
7	52.5	35.5	506	1	MATK_RHOFR
8	52.5	35.5	506	1	MATK_RHOTS
9	52	35.1	370	1	AROG_YEAST
10	51	34.5	232	1	AP3_ARATH
11	51	34.5	851	1	CE05_MOUSE
12	50	33.8	1378	1	RPOB_CAMJE
13	49	33.1	453	1	RMOG_PSEAE
14	48	32.4	205	1	RAS3_RHIRA
15	48	32.4	220	1	6POL_THEMA
16	48	32.4	519	1	THPC_SOLTU
17	48	32.4	526	1	THRC_ARATH
18	47.5	32.1	334	1	FMR1_CALPA
19	47.5	32.1	507	1	MATK_LOTFR
20	47	31.8	198	1	BIM_HUMAN
21	47	31.8	287	1	PRFA_POLPE
22	46.5	31.4	429	1	FMR2_ATEL
23	46.5	31.4	435	1	FMR1_ATEL
24	46	31.1	946	1	ITH2_HUMAN
25	45.5	30.7	1014	1	UVRA_STRCO
26	45	30.4	273	1	PSBO_ANASP
27	45	30.4	328	1	SNF4_KLULA
28	45	30.4	590	1	DCOA_SALTY
29	45	30.4	595	1	DCOA_KLEPN
30	45	30.4	653	1	HTZ2_HUMAN
31	45	30.4	865	1	ENY_STVAT
32	45	30.4	915	1	CE05_HUMAN
33	45	30.4	1535	1	LM11_CAEEL

RESULT 1

ID	BAD_MOUSE	STANDARD;	PRT;	204 AA.
AC	Q61337;			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	28-FEB-2003 (Rel. 41, Last annotation update)			
DE	Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-XL/Bcl-2 associated death promoter).			
DE	BAD OR BCL6.			
GN	Mus musculus (Mouse).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
OX	NCBI_TaxID=10090;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Brain, and Thymus;			
RX	MEDLINE=95136361; PubMed=7834748;			
RA	Yang E., Zha J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;			
RT	"Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and promotes cell death.";			
RL	Cell 80:285-291(1995).			
RN	[2]			
RP	PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.			
RX	MEDLINE=98022383; PubMed=9381178;			
RA	Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;			
RT	"Interleukin-3-induced phosphorylation of BAD through the protein kinase Akt.";			
RL	Science 278:687-689(1997).			
RN	[3]			
RP	MUTAGENESIS OF SERINE RESIDUES.			
RX	MEDLINE=20403302; PubMed=10949026;			
RA	Datta S.R., Katsov A., Hu L., Petros A., Yaffe M.B., Greenberg M.E.;			
RT	"I4-3-3 proteins and survival kinases cooperate to inactivate BAD by BH3 domain phosphorylation.";			
RL	Mol. Cell 6:41-51(2000).			
CC	-I- FUNCTION: Promotes cell death. Successfully competes for the binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level of heterodimerization of these proteins with BAX. Can reverse the death repressor activity of Bcl-x(L), but not that of Bcl-2. Appears to act as a link between growth factor receptor signaling and the apoptotic pathways.			
CC	-I- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity). The Ser-112/Ser-136 phosphorylated form binds I4-3-3 proteins.			
CC	-I- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon phosphorylation, locates to the cytoplasm.			
CC	-I- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND BAX for their pro-apoptotic activity and for their interaction with anti-apoptotic members of the Bcl-2 family.			
CC	-I- PTM: Phosphorylated on Ser-112 in response to survival stimuli. Subsequent phosphorylation on Ser-136 promotes heterodimerization with I4-3-3 proteins. This interaction then facilitates the phosphorylation at Ser-155, a site within the BH3 domain, leading to the release of Bcl-x(L) and the promotion of cell survival.			

34	45	30.4	5596	1	MDM1_HUMAN
35	44.5	30.1	506	1	MATK_GAUFR
36	44.5	30.1	512	1	MATK_LILFS
37	44.5	30.1	907	1	NUOG_ECOLI
38	44.5	30.1	907	1	NUOG_SALTY
39	44	29.7	196	1	BIM_MOUSE
40	44	29.7	196	1	BIM_RAT
41	44	29.7	262	1	END8_ECO57
42	44	29.7	262	1	END8_ECOLI
43	44	29.7	262	1	END8_SALTY
44	44	29.7	262	1	END8_SALTY
45	44	29.7	629	1	SYM_THEMA

ALIGNMENTS

Query Match 93.2%; Score 138; DB 1; Length 205;
 Best Local Similarity 100.0%; Pred. No. 2.5e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRLRRMSDEFGSKGL 27
 Db 141 NLWAAQRYGRLRRMSDEFGSKGL 166

RESULT 3
 ID BAD_HUMAN STANDARD; PRT; 168 AA.
 AC Q92934; O14803;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-
 DE XL/Bcl-2 associated death promoter) (BCL2-like 8 protein).
 GN BAD OR BOC6 OR BCL2L8.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Yin D.X., Li Z., Huang B., Chen S., Zhou H.;
 RT "A human protein that interacts with Bcl-2 and have homology to mouse
 RT BAD.";
 RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.
 RX MEDLINE=97083574; PubMed=8929532;
 RA Wang H.-G., Rapp U.R., Reed J.C.;
 RT "Bcl-2 targets the protein Kinase Raf-1 to mitochondria.";
 RL Cell 87:629-638(1996).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Takayama S., Reed J.C.;
 RL Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A., AND DIMERIZATION.
 RC TISSUE=Bone marrow;
 RX MEDLINE=98049554; PubMed=9388232;
 RA Ohtsue S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,
 RA Chang S., Weeks S., Fritz L.C., Oltersdorf T.;
 RT "Dimerization properties of human BAD.";
 RL J. Biol. Chem. 272:30866-30872(1997).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lung;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg K.B., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max A.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Ustin T.B., Toshlyuk S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fhney J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalls D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [6]

RP STRUCTURE BY NMR OF 103-127.
 RX MEDLINE=21073561; PubMed=11206074;
 RA Petros A.M., Nettelsheim D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
 RA Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
 RA Fesik S.W.;
 RT "Rationale for Bcl-xL/Bad peptide complex formation from structure,
 RT mutagenesis, and biophysical studies.";
 RL Protein Sci. 9:2528-2534(2000).
 CC -!- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 CC similarity). Appears to act as a link between growth factor
 CC receptor signaling and the apoptotic pathways.
 CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 CC The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
 CC similarity).
 CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm.
 CC -!- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
 CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -!- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-99 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-118, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-99 is the major site of AKT/PKB phosphorylation. Ser-118 the
 CC major site of protein kinase A (CAPK) phosphorylation (by
 CC similarity).
 CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
 CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -!- CAUTION: Ref.1 sequence differs from that shown due to frameshifts
 CC in position 64 and 91.
 CC -----
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 CC or send an email to license@sib-sib.ch).
 CC -----
 DR EMBL; U66879; AAB36516.1; ALT_FRAME.
 DR EMBL; AF021792; AAB72092.1; -.
 DR EMBL; AF031523; AAB88124.1; -.
 DR EMBL; BC001901; AAB01901.1; -.
 DR PDB; 1G57; 07-FEB-01.
 DR Genew; HGNC:936; BAD.
 DR MIM; 603167; -.
 DR GO; GO:0005737; C:cytoplasm; NAS.
 DR GO; GO:0005741; C:mitochondrial outer membrane; NAS.
 DR GO; GO:0005515; F:protein binding activity; NAS.
 DR GO; GO:0008632; P:apoptotic program; TAS.
 DR GO; GO:0008911; P:induction of apoptosis; NAS.
 DR InterPro; IPR0000712; Bcl2_BH.
 DR PROSITE; PS01259; BH3; FALSE_NEG.
 KW Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
 FT DOMAIN 110 124
 FT MOD_RES 75 75
 FT MOD_RES 99 99
 FT MOD_RES 118 118
 FT MOD_RES 107 107
 FT VARIANT 106 121
 FT HELIX 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
 FT SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
 SQ

Query Match

77.0%; Score 114; DB 1; Length 168;

Best Local Similarity 91.7%; Pred. No. 7.6e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 NLWAAQRYGRLRRMSDFEGSFK 25
|||||
Db 103 NLWAAQRYGRLRRMSDFVDSFK 126

RESULT 4
ID ITH2_MESAU STANDARD; PRT; 946 AA.
AC P37279;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Inter-alpha-trypsin inhibitor heavy chain H2 precursor (ITI heavy
chain H2) (inter-alpha-inhibitor heavy chain 2) (H2).
GN ITH2.
OS Mesocricetus auratus (Golden hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;
OC Mesocricetus.
OX NCBI_TaxID=10036;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=97420688; PubMed=9276673;
RA Nakatani T., Suzuki Y., Yamamoto T., Sinohara H.;
RT "Molecular cloning and sequencing of cDNAs encoding three heavy-chain
precursors of the inter-alpha-trypsin inhibitor in Syrian hamster:
RT implications for the evolution of the inter-alpha-trypsin inhibitor
heavy chain family.";
RL J. Biochem. 122:71-82(1997).
RN [2]
RP SEQUENCE OF 55-64; 140-146; 151-156; 424-447; 500-528 AND 577-605,
AND SUBUNITS.
RC TISSUE=Plasma;
RX MEDLINE=97018241; PubMed=8864857;
RA Yamamoto T., Yamamoto K., Sinohara H.;
RT "Inter-alpha-trypsin inhibitor and its related proteins in Syrian
hamster urine and plasma.";
RL J. Biochem. 120:145-152(1996).
CC -!- FUNCTION: MAY ACT AS A CARRIER OF HYALURONAN IN SERUM OR AS A
CC BINDING PROTEIN BETWEEN HYALURONAN AND OTHER MATRIX PROTEIN,
CC INCLUDING THOSE ON CELL SURFACES IN TISSUES TO REGULATE THE
CC LOCALIZATION, SYNTHESIS AND DEGRADATION OF HYALURONAN WHICH ARE
CC ESSENTIAL TO CELLS UNDERGOING BIOLOGICAL PROCESSES (BY
CC SIMILARITY).
CC -!- SUBUNIT: I-ALPHA-I PLASMA PROTEASE INHIBITORS ARE ASSEMBLED FROM
CC ONE OR TWO HEAVY CHAINS (H1, H2 OR H3) AND ONE LIGHT CHAIN,
CC BIKUNIN. INTER-ALPHA-INHIBITOR (I-ALPHA-I) IS COMPOSED OF H1, H2
CC AND BIKUNIN, INTER-ALPHA-LIKE INHIBITOR (I-ALPHA-LI) OF H2 AND
CC BIKUNIN, AND PRE-ALPHA-INHIBITOR (P-ALPHA-I) OF H3 AND BIKUNIN.
CC -!- PFM: HEAVY CHAINS ARE INTERLINKED WITH BIKUNIN VIA A CHONDROITIN
CC 4-SULFATE BRIDGE TO THE THEIR C-TERMINAL ASPARTATE (BY
CC SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE ITH FAMILY.
CC -!- SIMILARITY: Contains 1 VWFA domain.
CC
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CC
CC EMBL; D89286; BAA13939.1; .
CC PIR; JC5575; JC5575.
CC InterPro; IPR006587; VIT.
CC DR InterPro; IPR002035; VWF_A.
CC Pfam; PF00092; vwa; 1.
CC SMART; SM00609; VIT; 1.

DR SMART: SM00327; VWFA; 1.
DR PROSITE; PS50234; VWFA; 1.
KW Serine protease inhibitor; Repeat; Signal; Multigene family;
KW Glycoprotein.
FT SIGNAL 1 18 POTENTIAL.
FT PROPEP 19 54 BY SIMILARITY.
FT CHAIN 55 702 INTER-ALPHA-TRYPsin INHIBITOR HEAVY CHAIN
H2.
FT PROPEP 703 946 BY SIMILARITY.
FT DOMAIN 308 468 VWFA.
FT CARBOHYD 118 118 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 263 263 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 445 445 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 578 578 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT BINDING 702 702 CHONDROITIN 4-SULFATE, CROSS-LINK SITE
(BY SIMILARITY).
FT CONFLICT 510 510 V -> Y (IN REF. 2).
FT CONFLICT 595 595 E -> I (IN REF. 2).
SQ SEQUENCE 946 AA; 106580 MW; CA8BF56458E7B2E CRC64;
Query Match 36.58; Score 54; DB 1; Length 946;
Best Local Similarity 34.6%; Pred. No. 4.3;
Matches 9; Conservative 5; Mismatches 12; Indels 0; Gaps 0;

Qy 2 NLWAAQRYGRLRRMSDFEGSFKGL 27
|.|. : | : ||| |.:
Db 212 NWVIVELQGMFLHVPDTFEGHFGV 237

RESULT 5
ID ITH2_MOUSE
AC Q61703; STANDARD; PRT; 946 AA.
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Inter-alpha-trypsin inhibitor heavy chain H2 precursor (ITI heavy
chain H2) (inter-alpha-inhibitor heavy chain 2).
GN ITH2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6N; TISSUE=Liver;
RX MEDLINE=95194326; PubMed=7534067;
RA Chan P., Risler J.-L., Raguenez G., Salier J.-P.;
RT "The three heavy-chain precursors for the inter-alpha-inhibitor
family in mouse: new members of the multicopper oxidase protein group
with differential transcription in liver and brain.";
RL Biochem. J. 306:505-512(1995).
CC -!- FUNCTION: MAY ACT AS A CARRIER OF HYALURONAN IN SERUM OR AS A
CC BINDING PROTEIN BETWEEN HYALURONAN AND OTHER MATRIX PROTEIN,
CC INCLUDING THOSE ON CELL SURFACES IN TISSUES TO REGULATE THE
CC LOCALIZATION, SYNTHESIS AND DEGRADATION OF HYALURONAN WHICH ARE
CC ESSENTIAL TO CELLS UNDERGOING BIOLOGICAL PROCESSES (BY
CC SIMILARITY).
CC -!- SUBUNIT: I-ALPHA-I PLASMA PROTEASE INHIBITORS ARE ASSEMBLED FROM
CC ONE OR TWO HEAVY CHAINS (H1, H2 OR H3) AND ONE LIGHT CHAIN,
CC BIKUNIN. INTER-ALPHA-INHIBITOR (I-ALPHA-I) IS COMPOSED OF H1, H2
CC AND BIKUNIN, INTER-ALPHA-LIKE INHIBITOR (I-ALPHA-LI) OF H2 AND
CC BIKUNIN, AND PRE-ALPHA-INHIBITOR (P-ALPHA-I) OF H3 AND BIKUNIN.
CC -!- TISSUE SPECIFICITY: EXPRESSED IN BOTH LIVER AND BRAIN.
CC -!- PFM: HEAVY CHAINS ARE INTERLINKED WITH BIKUNIN VIA A CHONDROITIN
CC 4-SULFATE BRIDGE TO THE THEIR C-TERMINAL ASPARTATE (BY
CC SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE ITH FAMILY.
CC -!- SIMILARITY: Contains 1 VWFA domain.
CC
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CC
CC EMBL; D89286; BAA13939.1; .
CC PIR; JC5575; JC5575.
CC InterPro; IPR006587; VIT.
CC DR InterPro; IPR002035; VWF_A.
CC Pfam; PF00092; vwa; 1.
CC SMART; SM00609; VIT; 1.

DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Maturase K (Intron maturase).
GN MATK.
OS Rhododendron tsusiophyllum.
OC Chloroplast.
OG Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
ON NCBI_TaxID=49629;
RX [1]
CC "Investigation of sectional relationships in the genus
RP Kurshige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RI Rhododendron (Ericaceae) based on matk sequences.";
RL Shokubutsu Kenkyu Zasshi 73:143-154(1998).
CC -!- FUNCTION: Probably assists in splicing chloroplast group II
CC introns (By similarity).
CC -!- SIMILARITY: BELONGS TO THE INTRON MATURASE FAMILY 2. MATK
CC SUBFAMILY.
CC -----
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CC -----
DR EMBL; AB012750; BAA25871.1; .
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
DR mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60569 MW; AEE12FF8809C223E CRC64;

Query Match 35.5%; Score 52.5; DB 1; Length 506;
Best Local Similarity 37.5%; Pred. No. 3.6;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRRMSDFEGSFK 25
I :||| :||| I :|||
Db 391 KPFWAALSDSDIIERFGRIYRNLSHYSGSLK 422

RESULT 9
AROG_YEAST STANDARD; PRT; 370 AA.
ID AC F32449;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Phospho-2-dehydro-3-deoxyheptonate aldolase, tyrosine-inhibited
DE (SC 4.1.2.15) (Phospho-2-keto-3-deoxyheptonate aldolase) (DAPH
DE synthetase) (3-deoxy-D-arabino-heptulosonate 7-phosphate synthase).
DN AR04 OR YBE1249C OR YBE1701.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes;
OX NCBI_TaxID=4932;
RX [1]
CC SEQUENCE FROM N.A.
RP MEDLINE=9225349; PubMed=1348717;
RX Kuenzler M., Paravicini G., Egli C., Inniger S., Braus G.H.;
FT Cloning, primary structure and regulation of the AR04 gene, encoding
FT the tyrosine-inhibited 3-deoxy-D-arabino-heptulosonate-7-phosphate
FT synthase from Saccharomyces cerevisiae.';
RL Gene 113:67-74(1992).
RN [2]
RP REVISIONS TO 205-207.

OC eurons II; Brassicales; Brassicaceae; Arabidopsis.
 OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Petal;
 RX MEDLINE=92154682; PubMed=1346756;
 RA Jack T., Brockman L.L., Meyerowitz E.M.;
 RT "The homeotic gene APETALA3 of Arabidopsis thaliana encodes a MADS
 box and is expressed in petals and stamens.";
 RL Cell 68:683-697(1992).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Landsberg erecta;
 RX MEDLINE=95036018; PubMed=7948893;
 RA Okamoto H., Yano A., Shiraishi H., Okada K., Shimura Y.;
 RT "Genetic complementation of a floral homeotic mutation, apetal3,
 with an Arabidopsis thaliana gene homologous to DEFICIENS of
 Antirrhinum majus.";
 RL Plant Mol. Biol. 26:465-472(1994).
 RN [3]
 RP SEQUENCE FROM N.A., AND VARIANTS.
 RC STRAIN=cv. Bla-1, cv. Bretagny, cv. Bs-1, cv. Bu-0, cv. Bu-2,
 cv. Chi-1, cv. Co-1, cv. Columbia, cv. Corsacalla-1, cv. Cvi-0,
 cv. Gr-3, cv. J1-1, cv. Kas-1, cv. Kent, cv. Landsberg erecta,
 cv. Li-3, cv. Li-8, and cv. Lisse;
 RX MEDLINE=99126449; PubMed=9927474;
 RA Prugnanan M.D., Suddith J.I.;
 RT "Molecular population genetics of floral homeotic loci: departures
 from the equilibrium-neutral model at the APETALA3 and PISTILLATA
 genes of Arabidopsis thaliana.";
 RL Genetics 151:839-848(1999).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=21016720; PubMed=11130713;
 RA Salanoubat M., Lemcke K., Rieger M., Ansoerge W., Unseld M.,
 Partmann B., Valle G., Bloeker H., Perez-Alonso M., Obermaier B.,
 Delsenly M., Boutry M., Grivell L.A., Mache R., Puigdomenech P.,
 De Simone V., Choisine N., Artiguenave F., Robert C., Brottier P.,
 Wincker P., Cattolico L., Weissbach J., Saurin W., Quettier F.,
 Schaefer M., Mueller-Auer S., Gabel C., Fuchs M., Benes V.,
 Wurmbach E., Drzonek H., Erfle H., Jordan N., Bangert S.,
 Wiedelmann R., Kranz H., Voss H., Holland R., Brandt P., Nyakatura G.,
 Verzi A., D'Angelo M., Pallavicini A., Toppo S., Simonati B.,
 Conrad A., Hornischer K., Kauer G., Loehner T.-H., Nordsiek G.,
 Reichelt J., Scharfe M., Schoen O., Bagues M., Ferol J., Climent J.,
 Navarro P., Collado C., Perez-Perez A., Ottenwaelder B., Duchemin D.,
 Cooke R., Laudie M., Berger-Llauro C., Purnelle B., Masuy D.,
 de Haan M., Maarse A.C., Alcaraz J.-P., Cottet A., Casacuberta E.,
 Monfort A., Argirou A., Flores M., Liguori R., Vitale D.,
 Mannhaupt G., Haase D., Schoof H., Rudd S., Zaccaria P., Mewes H.-W.,
 Mayer K.F.X., Kaul S., Town C.D., Koo H.L., Tallon L.J., Jenkins J.,
 Rooney T., Kirzo M., Walts A., Otterback T., Fujii C.Y., Shea T.P.,
 Creasy T.H., Haas B., Maiti R., Wu D., Peterson J., Van Aken S.,
 Pai G., Miltscher J., Sellers P., Gill J.E., Feldblum T.V.,
 Pruss D., Lin X., Nierman W.C., Salzberg S.L., White O., Venter J.C.,
 Fraser C.M., Kaneko T., Nakamura Y., Sato S., Kato T., Asamizu E.,
 Sasamoto S., Kimura T., Idesawa K., Kawashima K., Kishida Y.,
 Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
 Nakayama S., Nakazaki N., Shinpo S., Takeuchi C., Wada T.,
 Watanabe A., Yamada M., Yasuda M., Tabata S.;
 RT "Sequence and analysis of chromosome 3 of the plant Arabidopsis
 thaliana.";
 RL Nature 408:820-822(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RA Brover V., Troukhan M., Alexandrov N., Lu Y.-P., Flavell R.,
 Feldmann K.;
 RT "Full-length cDNA from Arabidopsis thaliana.";
 RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RA Shinozaki K., Davis R.W., Ecker J.R., Theologis A.;
 RT "RIKEN Arabidopsis full length cDNA clones (RAFLs) sequenced by the
 SSP consortium (Salk/Stanford/PGEC).";
 RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
 RN [7]
 RP SEQUENCE OF 36-128 FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=99311297; PubMed=10382288;
 RA Brunel D., Froger N., Pelletier G.;
 RT "Development of amplified consensus genetic markers (ACGM) in Brassica
 napus from Arabidopsis thaliana sequences of known biological
 function.";
 RL Genome 42:387-402(1999).
 RN [8]
 RP FUNCTION.
 RX PubMed=8565821;
 RA Krizek B.A., Meyerowitz E.M.;
 RT "The Arabidopsis homeotic genes APETALA3 and PISTILLATA are sufficient
 to provide the B class organ identity function.";
 RL Development 122:11-22(1996).
 RN [9]
 RP CHARACTERIZATION.
 RX PubMed=8643482;
 RA Riechmann J.L., Krizek B.A., Meyerowitz E.M.;
 RT "Dimerization specificity of Arabidopsis MADS domain homeotic proteins
 APETALA1, APETALA3, PISTILLATA, and AGAMOUS.";
 RL Proc. Natl. Acad. Sci. U.S.A. 93:4793-4798(1996).
 RN [10]
 RP GENETIC REGULATION.
 RX PubMed=11283333;
 RA Ng M., Yanofsky M.F.;
 RT "Activation of the Arabidopsis B class homeotic genes by APETALA1.";
 RL Plant Cell 13:739-753(2001).
 RN [11]
 RP CHARACTERIZATION.
 RX PubMed=11206550;
 RA Honma T., Goto K.;
 RT "Complexes of MADS-box proteins are sufficient to convert leaves into
 floral organs.";
 RL Nature 409:525-529(2001).
 CC -I- FUNCTION: Probable transcription factor involved in the genetic
 control of flower development. Is required for normal development
 of petals and stamens in the wild-type flower. Forms an
 heterodimer with PISTILLATA that is required for autoregulation of
 both AP3 and PI genes. AP3/PI heterodimer interacts with APETALA1
 or SEPALLATA3 to form a ternary complex that could be responsible
 for the regulation of the genes involved in the flower
 development.
 CC -I- SUBUNIT: Forms an heterodimer with PISTILLATA, capable of binding
 to CARG-box sequences. AP3/PI heterodimer binds AP1 or SEP3 to
 form complexes.
 CC -I- SUBCELLULAR LOCATION: Nuclear.
 CC -I- TISSUE SPECIFICITY: Expressed in petals and stamens.
 CC -I- INDUCTION: Positively regulated by the meristem identity proteins
 APETALA1 and LEAFY with the cooperation of UFO.
 CC -I- MISCELLANEOUS: Mutations in AP3 cause transformation of petals
 into sepals and stamina into carpels.
 CC -I- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
 FACTORS.
 CC -I- SIMILARITY: Contains 1 K-box dimerization domain.
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 CC EMBL; M86357; AAA32740.1; -;
 DR EMBL; D21125; BAB04665.1; -;
 DR EMBL; AF115798; AAD51887.1; -;
 DR EMBL; AF115799; AAD51888.1; -;

DR	EMBL; AF115800;	AAD51889.1;	-
DR	EMBL; AF115801;	AAD51890.1;	-
DR	EMBL; AF115802;	AAD51891.1;	-
DR	EMBL; AF115803;	AAD51892.1;	-
DR	EMBL; AF115804;	AAD51893.1;	-
DR	EMBL; AF115805;	AAD51894.1;	-
DR	EMBL; AF115806;	AAD51895.1;	-
DR	EMBL; AF115807;	AAD51896.1;	-
DR	EMBL; AF115808;	AAD51897.1;	-
DR	EMBL; AF115809;	AAD51898.1;	-
DR	EMBL; AF115810;	AAD51899.1;	-
DR	EMBL; AF115811;	AAD51900.1;	-
DR	EMBL; AF115812;	AAD51901.1;	-
DR	EMBL; AF115813;	AAD51902.1;	-
DR	EMBL; AF115814;	AAD51903.1;	-
DR	EMBL; ALI32971;	CAB81799.1;	-
DR	EMBL; AY087369;	AAM64919.1;	-
DR	EMBL; AY070397;	AAL49893.1;	-
DR	EMBL; AY142590;	AAN13159.1;	-
DR	EMBL; AF056541;	AAD41557.1;	-
DR	PIR; A42095;	A42095.	
DR	HSPF; P11746;	IYMW.	
DR	TRANSFAC; TOL776;	--	
DR	InterPro; IPR002487;	TF_Kbox.	
DR	InterPro; IPR002100;	TF_MADSbox.	
DR	Pfam; PF01486;	K-box; 1.	
DR	Pfam; PF00319;	SRF-TF; 1.	
DR	PRINTS; PF00404;	MADSDOMAIN.	
DR	SMART; SM00432;	MADS; 1.	
DR	PROSITE; PS00350;	MADS_BOX_1; 1.	
DR	PROSITE; PS00066;	MADS_BOX_2; 1.	
KW	Flowering; transcription regulation;		
KW	Nuclear protein; DNA-binding; Coiled coil;		
KW	Polymorphism.		
FT	DOMAIN	3	57
FT	DOMAIN	93	165
FT	DOMAIN	75	164
FT	VARIANT	31	31
FT	VARIANT	47	47
FT	VARIANT	61	61
FT	VARIANT	73	73
FT	VARIANT	109	109
FT	VARIANT	115	115
Query Match		34.5%;	Score 51; DB 1; Length 232;
Best Local Similarity		44.4%;	Pred. No. 2.6;
Matches 12; Conservative		3; Mismatches	4; Indels 8; Gaps 1;
QY	7 QRYG-----RELRRMSDEFGSEFK 25		
		: :	:
Dd	107 QRLGCLDLDLDELRLLEDEMENIFK 133		
RESULT 11			
CE05_MOUSE	ID CE05_MOUSE	STANDARD;	PRT; 851 AA.
AC	Q8KZH3;		
DT	15-SEP-2003 (Rel. 42, Created)		
DT	15-SEP-2003 (Rel. 42, Last sequence update)		
DT	15-SEP-2003 (Rel. 42, Last annotation update)		
DE	Protein C5orf5 homolog.		
GN	C5ORF5.		
OS	Mus musculus (Mouse).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
OX	NCBI_TaxID=10090;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RC	STRAIN=FVB/N;		
EX	MEDLINE=22388257; PubMed=12477932;		
RA	Strausberg R.B., Feingold E.A., Grouse L.H., Derge J.G.,		
RA	Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,		
RA	Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,		

RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
RA Garber R.L., Goltry L., Tolentino E., Westbrook-Wadman S., Yuan Y.,
RA Brody R.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.,
RT "Complete genome sequence of *Pseudomonas aeruginosa* PA01, an
RT opportunistic pathogen.",
RL Nature 406:959-964(2000).
CC -!- FUNCTION: Involved in DNA recombination (By similarity).
CC -!- SIMILARITY: BELONGS TO THE RMC FAMILY.
CC -----
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CC -----
CC EMBL; AE004535; AAG04420.1; -
CC PIR; E83517; E83517.
CC InterPro: IPR003798; DUF195.
CC Pfam: PF02646; RmcC; 1.
CC DNA recombination; Coiled coil; Complete proteome.
CC FT DOMAIN 16 COILED COIL (POTENTIAL).
CC KW SEQUENCE 453 AA; 51539 MW; 1E7EA97E82EC5E4B CRC64;
CC -----
CC Query Match 33.1%; Score 49; DB 1; Length 453;
CC Best Local Similarity 55.6%; Pred. No. 11;
CC Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;
CC -----
CC QY 4 WAAQRYGR--ELRRMSDE 19
CC ||::: || ||::: ||
CC Db 65 WASERGREELRLASE 82
CC -----
CC RESULT 14
CC RAS3_RHIRA
CC ID RAS3_RHIRA STANDARD; PRT; 205 AA.
CC AC P22280;
CC DT 01-AUG-1991 (Rel. 19, Created)
CC DT 01-AUG-1991 (Rel. 19, Last sequence update)
CC DT 28-FEB-2003 (Rel. 41, Last annotation update)
CC DE Ras-like protein 3.
CC GN RAS3
CC OS Rhizomucor racemosus (Mucor circinelloides f. lusitanicus).
CC Eukaryota; Fungi; Zygomycota; Zygomycetes; Mucorales; Mucoraceae;
CC Mucor.
CC OX NCBI_TaxID=4841;
CC [1]
CC SEQUENCE FROM N.A.
CC RC STRAIN=ATCC 1216B;
CC RX MEDLINE=91061774; PubMed=1701021;
CC RA Casale W.L., McConnell D.G., Wang S.-Y., Lee Y.-J., Linz J.E.;
CC "Expression of a gene family in the dimorphic fungus *Mucor racemosus*
CC which exhibits striking similarity to human ras genes.";
CC Mol. Cell. Biol. 10:6654-6663(1990).
CC -!- ENZYME REGULATION: ALTERNATE BETWEEN AN INACTIVE FORM BOUND TO GDP
CC AND AN ACTIVE FORM BOUND TO GTP. ACTIVATED BY A GUANINE
CC NUCLEOTIDE-EXCHANGE FACTOR (GEF) AND INACTIVATED BY A GTPASE-
CC ACTIVATING PROTEIN (GAP).
CC -!- SUBCELLULAR LOCATION: Plasma membrane.
CC -!- DEVELOPMENTAL STAGE: IN SPORULATING MYCELIUM AND MUCH LESS IN
CC GERMLING AND YEAST.
CC -!- SIMILARITY: BELONGS TO THE SMALL GTPASE SUPERFAMILY. RAS FAMILY.
CC -----
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CC -----

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CC DR EMBL: M55177; AAA83379.1; -
DR PIR: C36365; C36365.
DR HSP: P01112; 1PLL.
DR InterPro: IPR0013577; GTPase_Ras.
DR InterPro: IPR001806; Ras_trnsfmg.
DR InterPro: IPR005225; Small_GTP.
DR Pfam: PF00071; ras; 1.
DR PRINTS; PRO0449; RASTRNSFRMG.
DR SMART; SM00173; RAS; 1.
DR TIGRfams; TIGR00231; small_gtp; 1.
KW GTP-binding; Prenylation; Lipoprotein.
FT NP_BIND 16 23 GTP (BY SIMILARITY).
FT NP_BIND 53 67 GTP (BY SIMILARITY).
FT NP_BIND 122 125 GTP (BY SIMILARITY).
FT DOMAIN 38 46 EFFECTOR REGION (PROBABLE).
FT LIPID 202 202 FARNESYL (BY SIMILARITY).
SQ SEQUENCE 205 AA; 23408 MW; DBF086466F090F50 CRC64;

Query Match 32.4%; Score 48; DB 1; Length 205;
Best Local Similarity 62.5%; Pred. No. 6.3;
Matches 10; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 11 RELRMSDEFEFGSKG 26
DB 168 REIRMKKEQGRSKG 183

RESULT 15
6PGL_THEME
ID 6PGL_THEME STANDARD; PRT; 220 AA.
AC Q9X0N6;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 6-phosphogluconolactonase (EC 3.1.1.31) (6PGL).
GN PGL OR DEVB OR TM1154.
OS Thermotoga maritima.
OC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
OX NCBI_Taxid=2336;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MSB8 / DSM 3109;
RX MEDLINE=99287316; PubMed=10360571;
RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
RA McDonald L., Utterback T.R., Malek J.A., Linner K.D., Garrett M.M.,
RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
RA "Evidence for lateral gene transfer between Archaea and Bacteria from
RA genome sequence of Thermotoga maritima.";
RL Nature 399:323-329(1999).
CC -!- FUNCTION: HYDROLYSIS OF 6-PHOSPHOGLUCONOLACTONE TO 6-
CC PHOSPHOGLUCONATE.
CC -!- CATALYTIC ACTIVITY: 6-phospho-D-glucono-1,5-lactone + H(2)O = 6-
CC phospho-D-gluconate.
CC -!- PATHWAY: Pentose phosphate pathway; second step.
CC -!- SIMILARITY: BELONGS TO THE GLUCOSAMINE/GALACTOSAMINE-6-PHOSPHATE
CC ISOMERASE FAMILY. 6-PHOSPHOGLUCONOLACTONASE SUBFAMILY.
CC -----
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CC -----
CC EMBL: AE001772; AAD36230.1; -.
DR PIR: F72289; F72289.
DR TIGR; TM1154; -.
DR InterPro: IPR006148; Gluc_gal_isom.
```

```
DR InterPro: IPR005900; Phosphogluconlac.
DR Pfam; PF01182; Glucosamine_iso; 1.
DR TIGRfams; TIGR01198; pgl; 1.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 220 AA; 25325 MW; 950FD07EE01E60C3 CRC64;

Query Match 32.4%; Score 48; DB 1; Length 220;
Best Local Similarity 34.8%; Pred. No. 6.8;
Matches 8; Conservative 8; Mismatches 7; Indels 0; Gaps 0;

QY 5 AAQRYGRELRRMSDEFEFGSKGL 27
DB 111 ACEKYEREINSATDQFDLAILGM 133

Search completed: September 15, 2003, 17:23:01
Job time : 6.6 secs
```

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 ; Search time 30.4 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-55
Perfect score: 148
Sequence: 1 KNLWAAQRYGRELRLMSDEFGSFKGLK 28

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5
Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SPTREMBL23:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phase:*
10: sp_plant:*
11: sp_protist:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_virus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	91	61.5	146	13 Q919N2	Q919N2 brachydanio
2	57	38.5	471	17 Q8ZY71	Q8ZY71 pyrobaculum
3	53	35.8	196	16 Q8VJS3	Q8VJS3 mycobacteri
4	53	35.8	223	16 Q10843	Q10843 mycobacteri
5	53	35.8	946	11 Q8K016	Q8K016 mus musculu
6	52.5	35.5	505	8 Q47148	Q47148 menziesia c
7	52.5	35.5	506	8 Q47149	Q47149 rhododendro
8	52.5	35.5	506	8 Q47171	Q47171 rhododendro
9	52.5	35.5	506	8 Q63960	Q63960 rhododendro
10	52.5	35.5	506	8 Q62982	Q62982 rhododendro
11	52.5	35.5	506	8 Q62975	Q62975 rhododendro
12	52.5	35.5	506	8 Q62972	Q62972 rhododendro
13	52.5	35.5	506	8 Q62989	Q62989 rhododendro
14	52.5	35.5	506	8 Q62978	Q62978 rhododendro
15	52.5	35.5	506	8 Q47155	Q47155 rhododendro
16	52.5	35.5	506	8 Q47152	Q47152 rhododendro

17	52.5	35.5	506	8	Q47173	Q47173 rhododendro
18	52.5	35.5	506	8	Q62990	Q62990 rhododendro
19	52.5	35.5	506	8	Q62974	Q62974 rhododendro
20	52.5	35.5	506	8	Q62993	Q62993 menziesia m
21	52.5	35.5	506	8	Q47170	Q47170 rhododendro
22	52.5	35.5	506	8	Q47174	Q47174 rhododendro
23	52.5	35.5	506	8	Q62983	Q62983 rhododendro
24	52.5	35.5	506	8	Q62980	Q62980 rhododendro
25	52.5	35.5	506	8	Q62981	Q62981 rhododendro
26	52.5	35.5	506	8	Q62977	Q62977 rhododendro
27	52.5	35.5	506	8	Q47168	Q47168 menziesia p
28	52.5	35.5	506	8	Q62988	Q62988 rhododendro
29	52.5	35.5	506	8	Q62973	Q62973 rhododendro
30	52.5	35.5	506	8	Q62992	Q62992 ledum palus
31	52.5	35.5	506	8	Q47175	Q47175 rhododendro
32	52.5	35.5	506	8	Q8HSP1	Q8HSP1 rhododendro
33	52.5	35.5	506	8	Q8HSP0	Q8HSP0 rhododendro
34	52.5	35.5	506	8	Q8HSP9	Q8HSP9 rhododendro
35	52.5	35.5	506	8	Q8HSP8	Q8HSP8 rhododendro
36	52.5	35.5	506	8	Q8HSP7	Q8HSP7 rhododendro
37	52.5	35.5	506	8	Q8HSP6	Q8HSP6 rhododendro
38	52.5	35.5	506	8	Q8HSP5	Q8HSP5 rhododendro
39	52.5	35.5	506	8	Q8HSP4	Q8HSP4 rhododendro
40	52.5	35.5	507	8	Q62985	Q62985 rhododendro
41	52.5	35.5	507	8	Q62986	Q62986 rhododendro
42	52.5	35.5	508	8	Q62979	Q62979 rhododendro
43	51.5	34.8	506	8	Q47153	Q47153 rhododendro
44	51.5	34.8	506	8	Q47160	Q47160 rhododendro
45	51.5	34.8	506	8	Q8HSP4	Q8HSP4 rhododendro

ALIGNMENTS

RESULT 1

Q919N2 PRELIMINARY; PRT; 146 AA.
AC Q919N2;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-DEC-2001 (TRENBLrel. 19, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Bad.
GN Bad.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20373792; PubMed=10917738;
RA Inohara N., Nunez G.;
RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish."
RL Cell Death Differ. 7:509-510(2000).
DR EMBL: AF231017; AAF66962.2; -;
DR HSSP: Q92934; 1G5J.
DR 2FIN: ZDB-GENE-000616-1; bad.
SQ SEQUENCE 146 AA; 16546 MW; 28A5650B5107ECB CRC64;

Query Match 61.5%; Score 91; DB 13; Length 146;

Best Local Similarity 61.5%; Pred. No. 2.le-05; Mismatches 4; Indels 0; Gaps 0;

Matches 16; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

Qy 3 LWAQAQRYGRELRLMSDEFGSFKGLK 28
||||:||||:||||:|:|

Db 89 LWAQKYGQQLRLMSDEFGDKGMKVK 114

RESULT 2

Q8ZY71 PRELIMINARY; PRT; 471 AA.
ID Q8ZY71
AC Q8ZY71;

```

DT 01-MAR-2002 (TREMBLrel. 20, Created)
DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)
DE 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Hypothetical protein PAE0922.
GN PAE0922.
OS Pyrobaculum aerophilum.
OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;
OC Thermoproteaceae; Pyrobaculum.
OX NCBI_TaxID=13773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IM2 / ATCC 51768 / DSM 7523;
RX PubMed=11792869;
RA Fitz-Gibbon S.T., Ladner H., Kim U.-J., Stetter K.O., Simon M.I.,
RA Miller J.H.;
RT "genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum
RT aerophilum.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989(2002).
DR EMBL; AE009793; AAL63125.1; -.
DR InterPro; IPR006638; Elp3.
DR InterPro; IPR000182; GCN5acetyltransf.
DR Pfam; PF00583; Acetyltransf; 1.
DR SMART; SM00729; Elp3; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 471 AA; 52952 MW; 3B1E36E8AEE2EF0A CRC64;

Query Match 38.5%; Score 57; DB 17; Length 471;
Best Local Similarity 44.0%; Pred. No. 7;
Matches 11; Conservative 3; Mismatches 11; Indels 0; Gaps 0;

QY 4 WAAQRYGRELRRMSDFESFKGLK 28
| ||||| :::: | | | |
Db 404 WQHSSGMGRELRLAEIAGRGALK 428

RESULT 3
Q8VJS3
ID Q8VJS3 PRELIMINARY; PRT; 196 AA.
AC Q8VJS3;
DT 01-MAR-2002 (TREMBLrel. 20, Created)
DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE IS1507, transposase.
GN MT2070.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W.;
RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB007038; AAK46348.1; -.
DR TIGR; MT2070; -.
DR InterPro; IPR003346; Transposase_20.
DR Pfam; PF02371; Transposase_20; 1.
SQ SEQUENCE 196 AA; 21349 MW; C145A8D836FD9C2D CRC64;

Query Match 35.8%; Score 53; DB 16; Length 196;
Best Local Similarity 58.8%; Pred. No. 10;
Matches 10; Conservative 1; Mismatches 6; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSD 18
| ||||| | : | | |
Db 134 NLWAAADRYNRAIARGHD 150

RESULT 4
Q10843
ID Q10843 PRELIMINARY; PRT; 223 AA.
AC Q10843;
DT 01-NOV-1998 (TREMBLrel. 08, Created)
DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
DT 01-MAR-2002 (TREMBLrel. 20, Last annotation update)
DE Hypothetical protein RV2014.
GN RV2014 OR MTCY39.03C.
OS Mycobacterium tuberculosis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
CC -!- SIMILARITY: TO M.PARATUBERCULOSIS IS900.
DR EMBL; Z74025; CAA98415.1; -.
DR Tuberculist; RV2014; -.
DR InterPro; IPR003346; Transposase_20.
DR Pfam; PF02371; Transposase_20; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 223 AA; 24132 MW; 70456750017FEF37 CRC64;

Query Match 35.8%; Score 53; DB 16; Length 223;
Best Local Similarity 58.8%; Pred. No. 12;
Matches 10; Conservative 1; Mismatches 6; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSD 18
| ||||| | : | | |
Db 165 NLWAAADRYNRAIARGHD 181

RESULT 5
Q8K016
ID Q8K016 PRELIMINARY; PRT; 946 AA.
AC Q8K016;
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Inter-alpha trypsin inhibitor, heavy chain 2.
GN ITIH2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Strausberg R.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC034341; AAH34341.1; -.
DR MGD; MGI:96619; Itih2.
DR InterPro; IPR006587; VIT.
DR InterPro; IPR002035; VWF_A.
DR SMART; SM00609; VIT; 1.
DR SMART; SM00327; VWA; 1.
DR PROSITE; PS50234; VWF_A; 1.
SQ SEQUENCE 946 AA; 105945 MW; 8B17DBA71B85BC5C CRC64;

```

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CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
CC EMBL: U61332; AAB93753.1; -.
DR InterPro; IPR000442; Intron_mature2.
DR InterPro; IPR002866; MatK.N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK.N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60502 MW; 0009EA88CD28549F CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps

QY 1 KNWAA-----QRYGRELRLMSDEFGSEFK 25
   | :||| :||| :| :| :| :|
Db 391 KPWAALSDSIIRFGRIYRLNLSYSGSLK 422

RESULT 8
O47171 PRELIMINARY; PRT; 506 AA.
ID O47171
AC O47171;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Ribosomal maturase (intron maturase) (Maturase K).
DE MATK.
GN GN
OS Rhododendron edgeworthii.
OC Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodorea; Rhododendron.
OC NCBI_TaxID=49162;
RN [1]
RP R2
RA Kron K.A.;
RL "Phylogenetics of Rhododendroideae (Ericaceae).";
CC Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
CC EMBL: U61354; AAB93748.1; -.
DR InterPro; IPR000442; Intron_mature2.
DR InterPro; IPR002866; MatK.N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK.N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60485 MW; 8A6353BFC5F4DC85 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps

QY 1 KNWAA-----QRYGRELRLMSDEFGSEFK 25
   | :||| :||| :| :| :| :|
Db 391 KPWAALSDSIIRFGRIYRLNLSYSGSLK 422

RESULT 9
O63950 PRELIMINARY; PRT; 506 AA.
ID O63950
AC O63950;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Ribosomal maturase (intron maturase) (Maturase K).
DE MATK OR YCF14.
GN GN
OS Rhododendron tashiroi, and

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OS Rhododendron farrerae.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75582, 75580;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012749; BAA25870.1; -.
DR EMBL; AB012745; BAA25866.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
DR mRNA processing; Chloroplast.
KW SEQUENCE 506 AA; 60389 MW; DE0C07AEE608B787 CRC64;
SQ SEQUENCE 506 AA; 60389 MW; DE0C07AEE608B787 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRRMSDEFGESFK 25
   | :||| | :||| | :|||
Db 391 KPWAALSDDIIERFGRIYRNLSHYSGSLK 422

RESULT 10
O62982 PRELIMINARY; PRT; 506 AA.
AC O62982;
DT 01-AUG-1998 (TREMELrel. 07, Created)
DT 01-AUG-1998 (TREMELrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMELrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
DE INTRONS (BY SIMILARITY).
DE -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
DE Ribosomal maturase (Intron maturase) (Maturase K).
DE MATK.
GN MATK.
OS Rhododendron nipponicum.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75577;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012739; BAA25860.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
DR mRNA processing; Chloroplast.
KW SEQUENCE 506 AA; 60419 MW; 1F95132CCF4F6B40 CRC64;
SQ SEQUENCE 506 AA; 60419 MW; 1F95132CCF4F6B40 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

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QY 1 KNLWAA-----QRYGRELRRMSDEFGESFK 25
   | :||| | :||| | :|||
Db 391 KPWAALSDDIIERFGRIYRNLSHYSGSLK 422

RESULT 11
O62975 PRELIMINARY; PRT; 506 AA.
ID O62975;
AC O62975;
DT 01-AUG-1998 (TREMELrel. 07, Created)
DT 01-AUG-1998 (TREMELrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMELrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
DE MATK.
GN MATK.
OS Rhododendron ponticum.
OC Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49628;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012732; BAA25853.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
DR mRNA processing; Chloroplast.
KW SEQUENCE 506 AA; 60449 MW; 21DFF700B071B5B8 CRC64;
SQ SEQUENCE 506 AA; 60449 MW; 21DFF700B071B5B8 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRRMSDEFGESFK 25
   | :||| | :||| | :|||
Db 391 KPWAALSDDIIERFGRIYRNLSHYSGSLK 422

RESULT 12
O62972 PRELIMINARY; PRT; 506 AA.
ID O62972;
AC O62972;
DT 01-AUG-1998 (TREMELrel. 07, Created)
DT 01-AUG-1998 (TREMELrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMELrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
DE MATK.
GN MATK.
OS Rhododendron ovatum.
OC Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49169;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).

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CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012729; BAA25850.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60493 MW; D230F5458C20FEF0 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRLRRMSDEFGSFK 25
Db 391 KPVWAALSDSDIIEFRGRIYRLNLSHYSGSLK 422

RESULT 13
O62989 PRELIMINARY; PRT; 506 AA.
AC O62989;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron indicum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75581;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matk Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012735; BAA25856.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60489 MW; 6D38A1D4D6FEC9BF CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRLRRMSDEFGSFK 25
Db 391 KPVWAALSDSDIIEFRGRIYRLNLSHYSGSLK 422

RESULT 14
O62978 PRELIMINARY; PRT; 506 AA.
AC O62978;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.

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OS Rhododendron canadense.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49465;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matk Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012735; BAA25856.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60350 MW; 5E832589ED64EA25 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRLRRMSDEFGSFK 25
Db 391 KPVWAALSDSDIIEFRGRIYRLNLSHYSGSLK 422

RESULT 15
O47155 PRELIMINARY; PRT; 506 AA.
AC O47155;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron hongkongense.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49165;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;
RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; U61338; AAB93751.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60406 MW; 4B5C675CE32218D8 CRC64;

Query Match 35.5%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 34;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRLRRMSDEFGSFK 25

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Db 391 KPVWAALSDIERFGRIRNLSHYYSGLK 422

Search completed: September 15, 2003, 17:25:50
Job time : 31.4 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:01 ; Search time 38.1857 Seconds
(without alignments)
112.231 Million cell updates/sec

Title: US-09-544-664-56

Perfect score: 143
Sequence: 1 KNWAAQRYGRLRMSEDFEGSKGL (27)

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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23: /SIDSI/gcgdata/geneseg/geneseg-emb1/AA2002.DAT.*
24: /SIDSI/gcgdata/geneseg/geneseg-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	143	100.0	21	AA1980	Bcl2 polypeptide B
2	143	100.0	28	AA1981	Bcl2 polypeptide B
3	138	96.5	26	AA1982	Bcl2 polypeptide B
4	138	96.5	26	AA1983	Bcl2 polypeptide B
5	138	96.5	21	AA1984	Bcl2 polypeptide B
6	138	96.5	162	AA1985	Shorter murine BAD
7	138	96.5	204	AA1986	bcl-x(L)/bcl-2 ass
8	138	96.5	204	AA1987	Murine BCL-XL/BCL-2
9	138	96.5	204	AA1988	Mutant BCL-XL/BCL-2

10	138	96.5	204	19	AAW61317	Mutant BCL-XL/BCL-2
11	138	96.5	204	19	AAW61318	Mutant BCL-XL/BCL-2
12	138	96.5	204	19	AAW58332	Murine BAD protein
13	138	96.5	204	22	AAW70369	Longer murine BAD
14	138	96.5	204	24	ABR39082	Murine BAD protein
15	138	96.5	567	22	AAU00220	Bad-DTR apoptosis
16	114	79.7	24	23	AAU78627	Human Bad peptide
17	114	79.7	25	23	ABP56161	PTC-interacting T
18	114	79.7	25	23	ABG78484	Mutant Bcl2 compet
19	114	79.7	25	23	ABG78493	Mutant Bcl2 compet
20	114	79.7	25	23	AAU78610	Human Bad peptide
21	114	79.7	25	23	AAU78620	Human Bad peptide
22	114	79.7	166	18	AAW32476	BBC6 protein for r
23	114	79.7	168	19	AAW55779	Human Bcl-xL/Bcl-2
24	114	79.7	168	21	AAI13512	Human cell prolif
25	114	79.7	168	22	AAI13512	Human BAD mutant a
26	114	79.7	168	22	AAI13512	Human BAD protein.
27	114	79.7	168	22	AAI13512	Amino acid sequenc
28	114	79.7	168	24	ABR39081	Human BAD protein
29	114	79.7	201	23	ABP41630	Human ovarian anti
30	113	79.0	23	17	AAK95166	bcl-x(L)/bcl-2 ass
31	111	77.6	25	23	ABG78490	Mutant Bcl2 compet
32	111	77.6	25	23	AAU78617	Human Bad peptide
33	110	76.9	25	23	ABG78488	Mutant Bcl2 compet
34	110	76.9	25	23	ABG78489	Mutant Bcl2 compet
35	110	76.9	25	23	AAU78615	Human Bad peptide
36	110	76.9	25	23	AAU78616	Human Bad peptide
37	109	76.2	25	23	AAU78628	Human Bad peptide
38	109	76.2	25	23	ABG78486	Mutant Bcl2 compet
39	109	76.2	25	23	ABG78492	Mutant Bcl2 compet
40	109	76.2	25	23	ABG78497	Mutant Bcl2 compet
41	109	76.2	25	23	AAU78612	Human Bad peptide
42	109	76.2	25	23	AAU78619	Human Bad peptide
43	109	76.2	25	23	AAU78624	Human Bad peptide
44	108	75.5	25	23	ABG78485	Mutant Bcl2 compet
45	108	75.5	25	23	AAU78611	Human Bad peptide

ALIGNMENTS

RESULT 1
AA1980
ID AAB37056 standard; peptide; 27 AA.
XX AAB37056;
AC AAB37056;
XX 28-FEB-2001 (first entry)
DT Bcl2 polypeptide BH3 domain peptide #56.
DE Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiatic; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.
XX Homo sapiens.
XX OS
XX PN WO2000059526-A1.
XX 12-OCT-2000.
XX 06-APR-2000; 2000WO-US09352.
XX 07-APR-1999; 99US-0128202.
XX (UIJE-) UNIV JEFFERSON THOMAS.
XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX WPI; 2000-679325/66.
DR

XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer
XX
XX Claim 18; Page 19; 74pp; English.
XX
CC The invention relates to a peptide conjugate having the formula:
CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl optionally
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
XX
XX Sequence 27 AA;
XX
Query Match 100.0%; Score 143; DB 21; Length 27;
Best Local Similarity 100.0%; Pred. No. 7.3e-15;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 KNLWAQRYGRELRRMSDEFGSFKGL 27
Db 1 KNLWAQRYGRELRRMSDEFGSFKGL 27
RESULT 2
AAB37055
ID AAB37055 standard; peptide; 28 AA.
XX AAB37055;
XX
XX 28-FEB-2001 (first entry)
XX
XX Bcl2 polypeptide BH3 domain peptide #55.
XX
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
XX cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
XX apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
XX colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
XX melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
XX stroke; myocardial infarction.
XX
XX Homo sapiens.
XX
XX W0200059526-A1.
XX
XX 12-OCT-2000.
XX
XX 06-APR-2000; 2000WO-US09352.
XX
XX 07-APR-1999; 99US-0128202.
XX
XX (UYJE-) UNIV JEFFERSON THOMAS.
XX

PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX WPI; 2000-679325/66.
XX
XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer
XX
XX Claim 18; Page 19; 74pp; English.
XX
CC The invention relates to a peptide conjugate having the formula:
CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
XX
XX Sequence 28 AA;
XX
Query Match 100.0%; Score 143; DB 21; Length 28;
Best Local Similarity 100.0%; Pred. No. 7.6e-15;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 KNLWAQRYGRELRRMSDEFGSFKGL 27
Db 1 KNLWAQRYGRELRRMSDEFGSFKGL 27
RESULT 3
AAB37001
ID AAB37001 standard; peptide; 26 AA.
XX AAB37001;
XX
XX 28-FEB-2001 (first entry)
XX
XX Bcl2 polypeptide BH3 domain peptide #1.
XX
XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
XX cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
XX apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
XX colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
XX melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
XX stroke; myocardial infarction.
XX
XX Homo sapiens.
XX
XX W0200059526-A1.
XX
XX 12-OCT-2000.
XX
XX 06-APR-2000; 2000WO-US09352.
XX
XX 07-APR-1999; 99US-0128202.
XX

XX PA (UYJE-) UNIV JEFFERSON THOMAS.
XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX DR WPI; 2000-679325/66.
XX PT New peptide conjugates for modulating apoptosis or for inhibiting B
XX PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
XX PT treating neurodegenerative disorders, stroke, or cancer
XX PS Claim 18; Page 17; 74pp; English.
XX CC The invention relates to a peptide conjugate having the formula:
XX CC (R-X)n-peptide where n = 1-10; X = C-O, when the R-X group is attached
XX CC to the N-terminus of the peptide, or a side chain of the peptide where
XX CC the functional group of the side chain is NH2 or OH; or X = O or NH,
XX CC when the R-X group is attached to the C-terminus of the peptide, or a
XX CC side chain of the peptide, where the side chain functional group is COOH
XX CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one
XX CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
XX CC monosubstituted with a 1-5C straight or branched chain alkyl group,
XX CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
XX CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
XX CC of the peptide portion of the conjugate. The peptides represent analogues
XX CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
XX CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
XX CC useful for modulating apoptosis in the cells of a subject, or for
XX CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
XX CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
XX CC function. In particular, the peptide conjugate is useful for treating a
XX CC subject afflicted with a cancer characterized by cancer cells that
XX CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
XX CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
XX CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
XX CC conjugate is also useful for treating disorders characterized by
XX CC increased apoptosis, e.g. neurodegenerative disorders, acquired
XX CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
XX SQ Sequence 26 AA;
Query Match 96.5%; Score 138; DB 21; Length 26;
Best Local Similarity 100.0%; Pred. No. 4.1e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRLRRMSDEFGSFKGL 27
Db 1 NLWAAQRYGRLRRMSDEFGSFKGL 26
RESULT 4
AAB37002
ID AAB37002 standard; peptide; 26 AA.
XX AC AAB37002;
XX DT 28-FEB-2001 (first entry)
XX DE Bcl2 polypeptide BH3 domain peptide #2.
XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
XX KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
XX KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
XX KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
XX KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
XX KW stroke; myocardial infarction.
XX OS Homo sapiens.
XX PN W0200059526-A1.
XX PD 12-OCT-2000.
XX PN

PF 06-APR-2000; 2000WO-US09352.
XX PR 07-APR-1999; 99US-0128202.
XX PA (UYJE-) UNIV JEFFERSON THOMAS.
XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX DR WPI; 2000-679325/66.
XX PT New peptide conjugates for modulating apoptosis or for inhibiting B
XX PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
XX PT treating neurodegenerative disorders, stroke, or cancer
XX PS Claim 18; Page 17; 74pp; English.
XX CC The invention relates to a peptide conjugate having the formula:
XX CC (R-X)n-peptide where n = 1-10; X = C-O, when the R-X group is attached
XX CC to the N-terminus of the peptide, or a side chain of the peptide where
XX CC the functional group of the side chain is NH2 or OH; or X = O or NH,
XX CC when the R-X group is attached to the C-terminus of the peptide, or a
XX CC side chain of the peptide, where the side chain functional group is COOH
XX CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one
XX CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
XX CC monosubstituted with a 1-5C straight or branched chain alkyl group,
XX CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
XX CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
XX CC of the peptide portion of the conjugate. The peptides represent analogues
XX CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
XX CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
XX CC useful for modulating apoptosis in the cells of a subject, or for
XX CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
XX CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
XX CC function. In particular, the peptide conjugate is useful for treating a
XX CC subject afflicted with a cancer characterized by cancer cells that
XX CC express Bcl-2. The cancer includes cancers, neuroblastoma, melanoma, or
XX CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
XX CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
XX CC conjugate is also useful for treating disorders characterized by
XX CC increased apoptosis, e.g. neurodegenerative disorders, acquired
XX CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
XX SQ Sequence 26 AA;
Query Match 96.5%; Score 138; DB 21; Length 26;
Best Local Similarity 100.0%; Pred. No. 4.1e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRLRRMSDEFGSFKGL 27
Db 1 NLWAAQRYGRLRRMSDEFGSFKGL 26
RESULT 5
AAB37003
ID AAB37003 standard; peptide; 27 AA.
XX AC AAB37003;
XX DT 28-FEB-2001 (first entry)
XX DE Bcl2 polypeptide BH3 domain peptide #3.
XX KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
XX KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
XX KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
XX KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
XX KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
XX KW stroke; myocardial infarction.
XX OS Homo sapiens.
XX PN W0200059526-A1.
XX PD

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XX PD 12-OCT-2000.
XX OS
XX OS Synthetic.
XX PN WO200110888-A1.
XX XX
XX PD 15-FEB-2001.
XX PF 30-MAY-2000; 2000WO-US11864.
XX PR 28-MAY-1999; 99US-0136783.
XX PA (UJJE-) UNIV JEFFERSON THOMAS.
XX PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
XX DR WPI; 2000-679325/66.
XX PT New peptide conjugates for modulating apoptosis or for inhibiting B
XX PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
XX PT treating neurodegenerative disorders, stroke, or cancer
XX PS Claim 18; Page 17; 74pp; English.
XX CC The invention relates to a peptide conjugate having the formula:
XX CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
XX CC to the N-terminus of the peptide, or a side chain of the peptide where
XX CC the functional group of the side chain is NH2 or OH; or X = O or NH,
XX CC when the R-X group is attached to the C-terminus of the peptide, or a
XX CC side chain of the peptide, where the side chain functional group is COOH
XX CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkenyl containing one
XX CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
XX CC monosubstituted with a 1-5C straight or branched chain alkyl group,
XX CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
XX CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
XX CC of the peptide portion of the conjugate. The peptides represent analogues
XX CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
XX CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
XX CC useful for modulating apoptosis in the cells of a subject, or for
XX CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
XX CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
XX CC function. In particular, the peptide conjugate is useful for treating a
XX CC subject afflicted with a cancer characterized by cancer cells that
XX CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
XX CC non-small lung, renal, or thyroid cancers, neuroblastoma, melanoma, or
XX CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
XX CC conjugate is also useful for treating disorders characterized by
XX CC increased apoptosis, e.g. neurodegenerative disorders, acquired
XX CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
XX SQ Sequence 27 AA;
XX CC
Query Match 96.5%; Score 138; DB 21; Length 27;
Best Local Similarity 100.0%; Pred. No. 4.3e-14;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRELRRMSDEFGSGKGL 27
DB 1 NLWAAQRYGRELRRMSDEFGSGKGL 26
RESULT 6
AAB70370
ID AAB70370 standard; protein; 162 AA.
XX AC AAB70370;
XX DT
XX DT 02-MAY-2001 (first entry)
XX DE Shorter murine BAD mutant amino acid sequence SEQ ID NO:3.
XX CC Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
XX CC immunostimulant; neuroprotective; neurotropic; antiischaemic; vulnery;
XX CC cytoskeletal; antiviral; antiarthritic; antiinflammatory; wound healing;
XX CC immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
XX CC immunodeficiency disease; neurodegenerative disease; viral infection;
XX CC ischaemic cell death; reperfusion cell death; arthritis; infertility;
XX CC lymphoproliferative condition; inflammation; autoimmune disease.

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XX OS Mus musculus.
XX OS Synthetic.
XX PN WO200110888-A1.
XX XX
XX PD 15-FEB-2001.
XX PF 30-MAY-2000; 2000WO-US11864.
XX PR 28-MAY-1999; 99US-0136783.
XX PA (APOP-) APOPTOSIS TECHNOLOGY INC.
XX PI Zhou X;
XX DR WPI; 2001-138734/14.
XX PT New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
XX PT useful for screening for candidate compounds which induce or inhibit
XX PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
XX PT Ser113 -
XX PS Claim 7; Page 148-149; 157pp; English.
XX CC The present invention describes an isolated or synthetic polypeptide
XX CC (I) comprising a less than full length amino acid sequence of a mutant
XX CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
XX CC fragment, which contains amino acid substitutions at Ser118 of a human
XX CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
XX CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
XX CC neurotropic, antiischaemic, vulnery, cytostatic, antiviral,
XX CC antiarthritic, antiinflammatory and immunosuppressive activities, and
XX CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
XX CC polynucleotides can be used for screening candidate compounds and drugs
XX CC for activity that promote cell survival or apoptosis. Other uses include
XX CC inducing or inhibiting apoptosis in a cell. Candidate compounds
XX CC identified and (mutant) BAD polypeptides are useful in treating
XX CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
XX CC death, reperfusion cell death, wound healing, cancer, viral infections,
XX CC lymphoproliferative conditions, arthritis, infertility, inflammation and
XX CC autoimmune diseases. The present sequence represents a specifically
XX CC claimed shorter murine BAD mutant amino acid sequence from the present
XX CC invention.
XX SQ Sequence 162 AA;
XX CC
Query Match 96.5%; Score 138; DB 22; Length 162;
Best Local Similarity 100.0%; Pred. No. 3.1e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 NLWAAQRYGRELRRMSDEFGSGKGL 27
DB 98 NLWAAQRYGRELRRMSDEFGSGKGL 123
RESULT 7
AAR95168
ID AAR95168 standard; Protein; 204 AA.
XX AC AAR95168;
XX DT
XX DT 06-JAN-1997 (first entry)
XX DE bcl-x(L)/bcl-2 associated death promoter protein.
XX CC Epitope; murine; bcl-x(L)/bcl-2 associated death promoter; Bad; stroke;
XX CC polypeptide; bcl-x; cell death; regulate; BH1; BH2; apoptotic cell death;
XX CC cytokine deprivation; IL-3 dependent cell line; immunodeficiency; AIDS;
XX CC neurodegenerative disease; senescence; ischaemia; neoplasia.
XX OS Mus musculus.
XX OS

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PH Key Location/Qualifiers
 FT Region 147..149
 FT /note= "BH1 conserved amino acids"
 FT Region 191..192
 FT /note= "BH2 conserved amino acids"
 FT Domain 38..61
 FT /note= "PEST sequence"
 FT Domain 111..130
 FT /note= "PEST sequence"
 XX WO9613614-A1.
 XX 09-MAY-1996.
 XX 31-OCT-1995; 95WO-US14246.
 XX 31-OCT-1994; 94US-0333565.
 XX (UNIW) UNIV WASHINGTON.
 XX Korsmeyer SJ;
 XX WPI; 1996-251465/25.
 XX N-PSDB; AAT29479.
 XX Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
 FT useful to treat neoplasia and apoptosis and to identify agents
 FT inhibiting its binding to bcl-2 or bcl-x(L) to form heteromultimers
 XX Claim 3; Fig 1; 130pp; English.
 XX This sequence represents the murine bcl-x(L)/bcl-2 associated death
 CC promoter (Bad) gene. Bad is a 22.1 kD protein which interacts with
 CC bcl-2 and bcl-x proteins and regulates cell death. It has homology
 CC to the bcl-2-related family clustered in the BHL and BH2 domain. Bad
 CC has been found to hybridise to bcl-x(L) and bcl-2 in yeast two-hybrid
 CC assays and in vivo in mammalian cells. Overexpressed Bad counters the
 CC death inhibitory activity of bcl-x(L), but is much less effective at
 CC countering the death inhibitory activity of bcl-2. Bad expression can
 CC accelerate apoptotic cell death induced by cytokine deprivation in an
 CC IL-3 dependent cell line expressing bcl-x(L), and its also counters the
 CC death repressor activity of bcl-x(L). Bad competes with Bax for binding
 CC to bcl-x(L). Bad may be used to identify agents which inhibit its
 CC binding to bcl-2 or bcl-x(L) to form heterodimers. Such agents may be
 CC used to treat neurodegenerative diseases, immunodeficiency diseases,
 CC e.g. AIDS, senescence or ischaemia.
 XX Sequence 204 AA;
 Query Match 96.5%; Score 138; DB 17; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRRMSDEFGSFKGL 27
 Db 140 NLWAAQRYGRELRRMSDEFGSFKGL 165
 RESULT 8
 AAW61315
 ID AAW61315 standard; Protein; 204 AA.
 XX AAW61315;
 XX 07-OCT-1998 (first entry)
 XX Murine BCL-XL/BCL-2 associated cell death regulator.
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX Mus sp.
 OS Synthetic.
 XX WO9817682-A1.

PN WO9817682-A1.
 XX 30-APR-1998.
 XX 17-OCT-1997; 97WO-US19175.
 XX 18-OCT-1996; 96US-0733505.
 XX (UNIW) UNIV WASHINGTON.
 XX Korsmeyer SJ;
 XX WPI; 1998-261422/23.
 XX N-PSDB; AAV27833.
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 FT useful for, e.g. treating reduced apoptosis such as in cancer or
 FT viral infection
 XX Claim 1; Fig 10; 95pp; English.
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence is the murine BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX Sequence 204 AA;
 Query Match 96.5%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRRMSDEFGSFKGL 27
 Db 140 NLWAAQRYGRELRRMSDEFGSFKGL 165
 RESULT 9
 AAW61316
 ID AAW61316 standard; Protein; 204 AA.
 XX AAW61316;
 XX 07-OCT-1998 (first entry)
 XX Mutant BCL-XL/BCL-2 associated cell death regulator #1.
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX Mus sp.
 OS Synthetic.
 XX WO9817682-A1.


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XX 30-APR-1998.
PD
XX
XX 17-OCT-1997; 97WO-US19175.
PF
XX
XX 18-OCT-1996; 96US-0733505.
PR
XX
XX (UNIW ) UNIV WASHINGTON.
PA
XX
XX Kormsmeier SJ;
PI
XX
XX WPI; 1998-261422/23.
DR
XX
XX N-PSDB; AAV27834.
DR
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
PT useful for, e.g. treating reduced apoptosis such as in cancer or
PT viral infection
XX
XX
XX Claim 7; Page 59; 95pp; English.
PS
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
CC death regulator) proteins, having an amino acid other than Ser at
CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
CC present sequence represents a mutant BAD protein. Also described are: (1)
CC fragments of mutant BAD protein able to decrease cell viability; (2)
CC fusion proteins of mutant BAD with a heterologous polypeptide that
CC increases intracellular delivery. Mutant BAD proteins are used to treat
CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
CC viral infection, lymphoproliferation, arthritis, infertility,
CC inflammation and autoimmune disease. Polynucleotide sequences encoding
CC mutant BAD proteins can be used similarly by gene therapy or to produce
CC transgenic animals for use as disease models or in drug screening. BAD
CC proteins phosphorylated at specified Ser are used to screen for enhancers
CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
CC aging or ischaemic cell death. The apoptotic status of cells is
CC determined by measuring relative amounts of phosphorylated and non-
CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
CC greater death-promoting activity than wild-type BAD which can become
CC phosphorylated on the specified Ser, forming a product that does not
CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
CC proteins in the cytosol, thus promoting cell survival. The mutants with
CC Ser substituted cannot bind 14-3-3.
XX
XX
XX Sequence 204 AA;
SQ
Query Match 96.5%; Score 138; DB 19; Length 204;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 NLWAAQRYGRLRRMSDEFGSKGL 27
Db 140 NLWAAQRYGRLRRMSDEFGSKGL 165
RESULT 10
AAW61317
ID AAW61317 standard; Protein; 204 AA.
XX
XX AAW61317;
AC
XX
XX 07-OCT-1998 (first entry)
DT
XX
XX Mutant BCL-XL/BCL-2 associated cell death regulator #2.
DE
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
KW serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
OS Synthetic.
XX
XX W09817682-A1.
PN
XX
XX 30-APR-1998.
PD

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PD
XX
XX 30-APR-1998.
XX
XX 17-OCT-1997; 97WO-US19175.
PF
XX
XX 18-OCT-1996; 96US-0733505.
PR
XX
XX (UNIW ) UNIV WASHINGTON.
PA
XX
XX Kormsmeier SJ;
PI
XX
XX WPI; 1998-261422/23.
DR
XX
XX N-PSDB; AAV27835.
DR
XX
XX New mutant BAD polypeptide with phosphorylatable serine replaced -
PT useful for, e.g. treating reduced apoptosis such as in cancer or
PT viral infection
XX
XX
XX Claim 7; Page 60; 95pp; English.
PS
XX
XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
CC death regulator) proteins, having an amino acid other than Ser at
CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
CC present sequence represents a mutant BAD protein. Also described are: (1)
CC fragments of mutant BAD protein able to decrease cell viability; (2)
CC fusion proteins of mutant BAD with a heterologous polypeptide that
CC increases intracellular delivery. Mutant BAD proteins are used to treat
CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
CC viral infection, lymphoproliferation, arthritis, infertility,
CC inflammation and autoimmune disease. Polynucleotide sequences encoding
CC mutant BAD proteins can be used similarly by gene therapy or to produce
CC transgenic animals for use as disease models or in drug screening. BAD
CC proteins phosphorylated at specified Ser are used to screen for enhancers
CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
CC aging or ischaemic cell death. The apoptotic status of cells is
CC determined by measuring relative amounts of phosphorylated and non-
CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
CC greater death-promoting activity than wild-type BAD which can become
CC phosphorylated on the specified Ser, forming a product that does not
CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
CC proteins in the cytosol, thus promoting cell survival. The mutants with
CC Ser substituted cannot bind 14-3-3.
XX
XX
XX Sequence 204 AA;
SQ
Query Match 96.5%; Score 138; DB 19; Length 204;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 NLWAAQRYGRLRRMSDEFGSKGL 27
Db 140 NLWAAQRYGRLRRMSDEFGSKGL 165
RESULT 11
AAW61318
ID AAW61318 standard; Protein; 204 AA.
XX
XX AAW61318;
AC
XX
XX 07-OCT-1998 (first entry)
DT
XX
XX Mutant BCL-XL/BCL-2 associated cell death regulator #3.
DE
XX
XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
KW serine substituted mutant; apoptosis; cancer; viral infection.
XX
XX Mus sp.
OS Synthetic.
XX
XX W09817682-A1.
PN
XX
XX 30-APR-1998.
PD

```

XX 17-OCT-1997; 97WO-US19175.
 PF 18-OCT-1996; 96US-0733505.
 XX (UNIW) UNIV WASHINGTON.
 XX Korsmeyer SJ;
 PI WPI; 1998-261422/23.
 DR N-PSDB; AAV27836.
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 XX Claim 7; Page 60-61; 95pp; English.
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX Ser substituted cannot bind 14-3-3.
 XX Sequence 204 AA;
 SQ Query Match 96.5%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSFKGL 27
 DB 140 NLWAAQRYGRELRLMSDEFGSFKGL 165
 RESULT 12
 AAWS5832
 ID AAWS5832 standard; protein; 204 AA.
 XX AC AAWS5832;
 XX 23-JUL-1998 (first entry)
 DT Murine BAD protein.
 DE BAD protein; Bcl-XL/Bcl-2 associated cell death regulator; 14-3-3;
 KW serine phosphorylation; post-translational modification; apoptosis;
 KW signal transduction regulator; phosphoserine phosphatase; senescence;
 KW immunodeficiency disease; neurodegenerative disease; infertility;
 KW cancer; viral infection; lymphoproliferative condition; arthritis;
 KW inflammation; autoimmune diseases.
 XX Mus sp.
 OS W09809643-A1.
 XX

XX 12-MAR-1998.
 PD 09-SEP-1997; 97WO-US15871.
 XX 09-SEP-1996; 96US-0707868.
 XX (UNIW) UNIV WASHINGTON.
 XX Korsmeyer SJ;
 PI WPI; 1998-207049/18.
 DR Serine-phosphorylated Bcl-X-L/Bcl-2 Associated cell Death regulator
 XX polypeptide - useful for modulation of apoptosis associated with,
 PT e.g. cancer and immunodeficiency diseases
 PT Claim 3; Fig 8; 61pp; English.
 PS This sequence represents a novel serine-phosphorylated protein, BAD
 CC (Bcl-XL/Bcl-2 associated cell death regulator). The serine residue is
 CC phosphorylated in a post-translational modification and allows binding
 CC to the 14-3-3 protein which is a signal transduction regulator.
 CC Modulators of phosphorylated BAD, which act through inhibition/activation
 CC of a phosphoserine phosphatase, are useful for preventing/treating
 CC increased/decreased apoptosis in a cell. The increased apoptosis may
 CC result from immunodeficiency diseases, senescence, neurodegenerative
 CC disease, ischaemic cell death, reperfusion cell death, infertility and
 CC wound-healing. Decreased apoptosis may result from cancer, viral
 CC infection, lymphoproliferative conditions, arthritis, infertility,
 CC inflammation and autoimmune diseases. Measuring the amount of
 CC phosphorylated compared to unphosphorylated BAD polypeptide and/or total
 CC BAD in a cell is useful for determining the apoptotic state of a cell.
 XX Sequence 204 AA;
 SQ Query Match 96.5%; Score 138; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDEFGSFKGL 27
 DB 140 NLWAAQRYGRELRLMSDEFGSFKGL 165
 RESULT 13
 AAB70369
 ID AAB70369 standard; protein; 204 AA.
 XX AC AAB70369;
 XX 02-MAY-2001 (first entry)
 DT Longer murine BAD mutant amino acid sequence SEQ ID NO:2.
 DE Bcl-XL/Bcl-2 associated cell death regulator; BAD; mutant; apoptosis;
 XX immunostimulant; neuroprotective; nootropic; antischaeamic; vulnary;
 KW cytosolic; antiviral; antiarthritic; antiinflammatory; wound healing;
 KW immunosuppressive; apoptosis inducer; apoptosis inhibitor; cancer;
 KW immunodeficiency disease; neurodegenerative disease; viral infection;
 KW ischaemic cell death; reperfusion cell death; arthritis; infertility;
 KW lymphoproliferative condition; inflammation; autoimmune disease.
 XX Mus musculus.
 OS Synthetic.
 XX W0200110888-A1.
 PN 15-FEB-2001.
 PD 30-MAY-2000; 2000WO-US11864.
 XX 28-MAY-1999; 99US-0136783.
 XX

XX PA (APOP-) APOPTOSIS TECHNOLOGY INC.
 XX PI Zhou X;
 XX WPI; 2001-138734/14.
 XX New mutant Bcl-XL/Bcl-2 Associated Cell Death Regulator polypeptide,
 PT useful for screening for candidate compounds which induce or inhibit
 PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 PT Ser113 -
 XX Claim 4; Page 148; 157pp; English.
 XX The present invention describes an isolated or synthetic polypeptide
 CC comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (I) has immunostimulant, neuroprotective,
 CC neurotropic, antischismic, vulnerary, cytostatic, antiviral, and
 CC antiarthritic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed longer murine BAD mutant amino acid sequence from the present
 CC invention.
 XX SQ Sequence 204 AA;
 Query Match 96.5%; Score 138; DB 22; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDFEGSFKGL 27
 DB 140 NLWAAQRYGRELRLMSDFEGSFKGL 165
 RESULT 14
 ABR39082
 ID ABR39082 standard; Protein; 204 AA.
 XX AC ABR39082;
 XX 10-MAY-2003 (first entry)
 DE Murine BAD protein SEQ ID NO.4.
 XX Murine; BAD; herpes simplex virus; HSV; US3; herpes virus; apoptosis;
 KW virucide; infection.
 XX OS Mus musculus.
 XX WO2003012049-A2.
 PN 13-FEB-2003.
 XX 31-JUL-2002; 2002WO-US241177.
 PF 31-JUL-2001; 2001US-308929P.
 XX (UYCH-) UNIV CHICAGO.
 PA Munger J, Roizman B;
 XX 2003-248168/24.
 DR WPI; 2003-248168/24.

DR N-PSDB; ABZ61201.
 XX Inducing apoptosis in a cell infected with herpes simplex virus, HSV,
 PT by administering to the cell, a composition comprising an agent that
 PT inhibits phosphorylation of pro-apoptotic polypeptide BAD by HSV US3 -
 XX Claim 15; Page 168; 192pp; English.
 XX The present invention describes a method (M1) for inducing apoptosis in
 CC a cell infected with herpes simplex virus (HSV), which comprises
 CC administering to the cell, a composition having an agent that inhibits
 CC phosphorylation of pro-apoptotic polypeptide BAD by HSV US3. Also
 CC described is a method (M2) for treating a patient infected with HSV, by
 CC administering to the patient, a composition comprising a peptide
 CC comprising a sequence of 4-100 continuous amino acids of a 168 residue
 CC amino acid sequence (see ABR39081), where the peptide comprises ser112,
 CC ser135, or ser155, or their combinations. BAD has virucide activity.
 CC M1 is useful for inducing apoptosis in a cell infected with HSV, where
 CC the cell is in a human. M2 is useful for treating a patient infected
 CC with HSV. The present sequence represents murine BAD, which is used in
 CC the exemplification of the present invention.
 XX SQ Sequence 204 AA;
 Query Match 96.5%; Score 138; DB 24; Length 204;
 Best Local Similarity 100.0%; Pred. No. 4e-13;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 NLWAAQRYGRELRLMSDFEGSFKGL 27
 DB 140 NLWAAQRYGRELRLMSDFEGSFKGL 165
 RESULT 15
 AAU00220
 ID AAU00220 standard; Protein; 567 AA.
 XX AC AAU00220;
 XX 31-MAY-2001 (first entry)
 DE Bad-DTRR apoptosis-modifying fusion protein.
 XX Mouse; Bad-DTRR; apoptosis; cancer; spinal muscular atrophy;
 KW diphtheria toxin receptor binding domain; DTR; neoplasm; tumour;
 KW hyper-proliferation; Alzheimer's disease; neurodegenerative disorder;
 KW transient ischaemic neuronal injury; stroke; spinal cord injury;
 KW Huntington's disease.
 XX Chimeric - Mus sp.
 OS Chimeric - Corynebacterium diphtheriae.
 OS Chimeric - Synthetic.
 XX Key Location/Qualifiers
 FT Region 3..12 /note= "10x histidine tag"
 FT WC200112661-A2.
 XX 22-FEB-2001.
 XX 15-AUG-2000; 2000WO-US22293.
 XX 16-AUG-1999; 99US-0149220.
 XX (HARD) HARVARD COLLEGE.
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX Youle RJ, Liu X, Collier RJ;
 XX WPI; 2001-218343/22.
 DR N-PSDB; AAS00248.

PT Novel fusion protein for modifying apoptosis in target cell and
 PT reducing apoptosis after transient ischaemic neuronal injury, has two
 PT domains which targets protein to a cell and modifies apoptotic response
 PT of cell -

XX
 PS Claim 4; Page 59-61; 65pp; English.

XX
 CC The sequence represents the amino acid sequence of Bad-DTRR apoptosis-
 CC modifying fusion protein comprising Bad gene sequence fused via a short
 CC linker to diphtheria toxin translocation domain (DTRR). The
 CC functional apoptosis-modifying fusion protein is capable of binding a
 CC target cell and integrating into or crossing a cellular membrane of the
 CC target cell.. The apoptosis-modifying fusion protein comprises at least
 CC two domains: the DTR domain, which targets the fusion protein to the
 CC target cell and the Bcl-XL domain, which modifies an apoptotic response
 CC of the target cell. The fusion protein is useful for modifying
 CC (inhibiting or enhancing) apoptosis in a target cell, such as neuron,
 CC lymphocyte, cancer, neoplasm, macrophage, epithelial, stem, tumour or
 CC hyper-proliferative cell or an adipocyte. It is also useful for reducing
 CC apoptosis in a subject after transient ischaemic neuronal injury,
 CC especially spinal cord injury. The fusion protein may be used to treat
 CC various diseases and injury conditions through inhibition or enhancement
 CC of apoptotic cellular response, including neurodegenerative disorders
 CC such as Alzheimer's disease, Huntington's disease, spinal muscular
 CC atrophy, stroke episodes and unregulated cell growth as in tumours and
 CC various cancers. The apoptosis-modifying fusion protein can be delivered
 CC effectively throughout the body and targeted to selective tissue and
 CC cells.

XX
 SQ Sequence 567 AA;

Query Match 96.5%; Score 138; DB 22; Length 567;
 Best Local Similarity 100.0%; Pred. No. 1.2e-12;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFGSKGL 27
 Db 161 NLWAAQRYGRELRLMSDEFGSKGL 186

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 Job time : 38.1857 secs

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:22:21 ; Search time 14.0786 Seconds
(without alignments)
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Title: US-09-544-664-56
Perfect score: 143
Sequence: 1 KNLWAAQRYGRELKMSDEFEFSFKGL 27

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents_AA.*

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- 2: /cgn2.6/prodata/1/iaa/5B.COMB.pep.*
- 3: /cgn2.6/prodata/1/iaa/6A.COMB.pep.*
- 4: /cgn2.6/prodata/1/iaa/6B.COMB.pep.*
- 5: /cgn2.6/prodata/1/iaa/PCRUS.COMB.pep.*
- 6: /cgn2.6/prodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	138	96.5	204	1 US-08-333-565-2	Sequence 2, Appli
2	138	96.5	204	2 US-08-661-479-2	Sequence 2, Appli
3	138	96.5	204	2 US-08-733-505A-1	Sequence 1, Appli
4	138	96.5	204	2 US-08-733-505A-12	Sequence 12, Appl
5	138	96.5	204	2 US-08-733-505A-13	Sequence 13, Appl
6	138	96.5	204	2 US-08-733-505A-14	Sequence 14, Appl
7	135	94.4	204	2 US-08-717-123-3	Sequence 3, Appli
8	135	94.4	204	4 US-09-375-257-3	Sequence 3, Appli
9	114	79.7	166	1 US-08-665-617-2	Sequence 2, Appli
10	114	79.7	168	2 US-08-717-123-2	Sequence 2, Appli
11	114	79.7	168	3 US-08-985-335-1	Sequence 1, Appli
12	114	79.7	168	3 US-08-985-335-7	Sequence 7, Appli
13	114	79.7	168	3 US-09-410-372-1	Sequence 1, Appli
14	114	79.7	168	3 US-09-410-372-7	Sequence 7, Appli
15	114	79.7	168	4 US-09-375-257-2	Sequence 2, Appli
16	113	79.0	23	1 US-08-333-565-10	Sequence 10, Appl
17	113	79.0	23	1 US-08-661-479-10	Sequence 10, Appl
18	102	71.3	59	2 US-08-733-505A-55	Sequence 55, Appl
19	102	71.3	59	2 US-08-733-505A-56	Sequence 56, Appl
20	102	71.3	59	2 US-08-733-505A-57	Sequence 57, Appl
21	102	71.3	59	2 US-08-733-505A-58	Sequence 58, Appl
22	86	60.1	16	1 US-08-333-565-26	Sequence 26, Appl
23	86	60.1	16	2 US-08-661-479-26	Sequence 26, Appl
24	61	42.7	11	2 US-08-733-505A-34	Sequence 34, Appl
25	61	42.7	11	2 US-08-706-741B-69	Sequence 69, Appl
26	61	42.7	11	2 US-08-924-695A-69	Sequence 69, Appl
27	51	35.7	66	2 US-08-867-087B-40	Sequence 40, Appl

28 48.5 33.9 904 4 US-09-328-352-4656 Sequence 4656, Ap
29 46 32.2 610 4 US-09-252-991A-19594 Sequence 19594, A
30 46 32.2 946 3 US-09-074-579-3 Sequence 3, Appli
31 46 32.2 946 3 US-09-388-774-3 Sequence 3, Appli
32 46 32.2 946 4 US-09-546-153-1 Sequence 1, Appli
33 45.5 31.8 906 4 US-09-252-991A-31458 Sequence 31458, A
34 45 31.5 229 4 US-09-252-991A-23807 Sequence 23807, A
35 45 31.5 303 4 US-09-328-352-5164 Sequence 5164, Ap
36 45 31.5 356 4 US-09-235-103-2 Sequence 2, Appli
37 45 31.5 356 4 US-09-235-103-4 Sequence 4, Appli
38 45 31.5 1064 4 US-09-252-991A-17508 Sequence 17508, A
39 44.5 31.1 903 4 US-09-252-991A-28775 Sequence 28775, A
40 44 30.8 125 4 US-09-328-352-7449 Sequence 7449, Ap
41 44 30.8 263 4 US-09-651-656-27 Sequence 27, Appl
42 44 30.8 263 4 US-09-650-855-27 Sequence 27, Appl
43 44 30.8 277 4 US-09-252-991A-28581 Sequence 28581, A
44 44 30.8 877 4 US-09-206-551-20 Sequence 20, Appl
45 44 30.8 1125 4 US-09-252-991A-18729 Sequence 18729, A

ALIGNMENTS

RESULT 1
US-08-333-565-2
; Sequence 2, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Kourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note="Deduced amino acid sequence
; OTHER INFORMATION: of mouse BAD."

US-08-333-565-2

Query Match 96.5%; Score 138; DB 1; Length 204;
Best Local Similarity 100.0%; Pred. No. 1e-13;
Matches 26; Conservative. 0; Mismatches 0; Indels 0; Gaps 0;

REFERENCE/DOCKET NUMBER: P-ID 1929
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
TOPOLOGY: linear
US-08-717-123-3

Query Match 94.4%; Score 135; DB 2; Length 204;
Best Local Similarity 96.2%; Pred. No. 2.9e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
|||||
Db 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 8
US-09-375-257-3
Sequence 3, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
FILE REFERENCE: 480140.428D1
CURRENT APPLICATION NUMBER: US/09/375,257
CURRENT FILING DATE: 1999-08-16
NUMBER OF SEQ ID NOS: 15
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 3
LENGTH: 204
TYPE: PRT
ORGANISM: Mus musculus
US-09-375-257-3

Query Match 94.4%; Score 135; DB 4; Length 204;
Best Local Similarity 96.2%; Pred. No. 2.9e-13;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
|||||
Db 140 NLWAAQRYGRELRRMTDEFGSKGL 165

RESULT 9
US-08-665-617-2
Sequence 2, Application US/08665617
Patent No. 5663316
GENERAL INFORMATION:
APPLICANT: Xudong, Yin
TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Saliwanchik & Saliwanchik
STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: Florida
COUNTRY: USA
ZIP: 32606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/665,617
FILING DATE:
CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:
NAME: Saliwanchik, David R.
REGISTRATION NUMBER: 31,794
REFERENCE/DOCKET NUMBER: CI-8
TELECOMMUNICATION INFORMATION:
TELEPHONE: (352) 375-8100
TELEFAX: (352) 372-5800
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 166 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-665-617-2

Query Match 79.7%; Score 114; DB 1; Length 166;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSK 25
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Db 101 NLWAAQRYGRELRRMSDEFGSK 124

RESULT 10
US-08-717-123-2
Sequence 2, Application US/08717123
Patent No. 5965703
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
TITLE OF INVENTION: Acids and Methods of Use
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell and Flores
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: United States
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/717,123
FILING DATE: 20-SEP-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-ID 1929
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-717-123-2

Query Match 79.7%; Score 114; DB 2; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSK 25
|||||
Db 103 NLWAAQRYGRELRRMSDEFGSK 126

```

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0855
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; US-08-985-335-7

Query Match          79.7%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2 NLWAAQRYGRELRRMSDEFEGSFK 25
      |||||
Db      103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 13
US-09-410-372-1
; Sequence 1, Application US/09410372
; Patent No. 6281334
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0855
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 358673
; US-08-985-335-1

Query Match          79.7%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2 NLWAAQRYGRELRRMSDEFEGSFK 25
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Db      103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 12
US-08-985-335-7
; Sequence 7, Application US/08985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9

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;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 08/985,335
;; FILING DATE:
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Billings, Lucy J.
;; REGISTRATION NUMBER: 36,749
;; REFERENCE/DOCKET NUMBER: PF-0421 US
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 650-845-0555
;; TELEFAX: 650-845-4166
;; INFORMATION FOR SEQ ID NO: 1:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 168 amino acids
;; TYPE: amino acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; IMMEDIATE SOURCE:
;; LIBRARY: SYNORAB01
;; CLONE: 358673
US-09-410-372-1
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Query Match 79.7%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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QY 2 NLWAAQRYGRELRRMSDEFEGSK 25
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126
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RESULT 14
US-09-410-372-7
; Sequence 7, Application US/09410372
; Patent No. 6281334
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/410,372
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/985,335
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-845-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
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;; TOPOLOGY: linear
;; IMMEDIATE SOURCE:
;; LIBRARY: GenBank
;; CLONE: 1683637
US-09-410-372-7
Query Match 79.7%; Score 114; DB 3; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFEGSK 25
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126

RESULT 15
US-09-375-257-2
; Sequence 2, Application US/09375257
; Patent No. 6504022
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oitersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.42801
; CURRENT APPLICATION NUMBER: US/09/375,257
; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-375-257-2
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Query Match 79.7%; Score 114; DB 4; Length 168;
Best Local Similarity 91.7%; Pred. No. 4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFEGSK 25
Db 103 NLWAAQRYGRELRRMSDEFVDSFK 126
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Job time : 15.0786 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model 1

Run on: September 15, 2003, 17:25:56 ; Search time 21.4071 Seconds
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Title: US-09-544-664-56

Perfect score: 143

Sequence: 1 KNLAAQRYGRLFRMSDEFEGSKGL 27

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Gapop 10.0 , Gapext 0.5

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Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:*

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- 12: /cgn2_6/ptodata/1/pubaa/US09_NEW_PUB.pep.*
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- 14: /cgn2_6/ptodata/1/pubaa/US10B_PUBCOMB.pep.*
- 15: /cgn2_6/ptodata/1/pubaa/US10C_PUBCOMB.pep.*
- 16: /cgn2_6/ptodata/1/pubaa/US10_NEW_PUB.pep.*
- 17: /cgn2_6/ptodata/1/pubaa/US60_NEW_PUB.pep.*
- 18: /cgn2_6/ptodata/1/pubaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	135	94.4	204	9	US-09-922-378-3
2	135	94.4	204	14	US-10-066-179-3
3	114	79.7	25	15	US-10-059-261-258
4	114	79.7	168	9	US-09-922-378-2
5	114	79.7	168	9	US-09-894-657-1
6	114	79.7	168	9	US-09-894-657-7
7	114	79.7	168	14	US-10-066-179-2
8	71	49.7	15	15	US-10-178-105A-147
9	51	35.7	682	12	US-10-238-075-1077
10	50	35.0	215	15	US-10-156-761-9145
11	47	32.9	35	15	US-10-092-750-1
12	47	32.9	138	15	US-10-092-750-241
13	46	32.2	946	9	US-09-828-423-3
14	44	30.8	272	15	US-10-156-761-11541
15	44	30.8	426	9	US-09-815-242-5704

16	44	30.8	699	14	US-10-008-355-8	Sequence 8, Appli
17	44	30.8	705	9	US-09-815-242-12463	Sequence 12463, A
18	44	30.8	712	14	US-10-008-355-2	Sequence 2, Appli
19	44	30.8	877	12	US-10-369-284-20	Sequence 20, Appli
20	43	30.1	213	9	US-03-843-846-18	Sequence 18, Appli
21	43	30.1	232	10	US-09-881-752A-238	Sequence 238, App
22	43	30.1	270	11	US-09-934-455A-162	Sequence 162, App
23	43	30.1	380	9	US-09-149-045-2	Sequence 2, Appli
24	43	30.1	380	15	US-10-166-359-2	Sequence 2, Appli
25	43	30.1	380	15	US-10-166-113-2	Sequence 2, Appli
26	43	30.1	380	15	US-10-166-357-2	Sequence 2, Appli
27	43	30.1	380	15	US-10-166-372-2	Sequence 2, Appli
28	43	30.1	380	15	US-10-184-722-3	Sequence 3, Appli
29	43	30.1	380	15	US-10-251-385-62	Sequence 62, Appli
30	43	30.1	380	15	US-10-251-385-198	Sequence 198, App
31	43	30.1	380	15	US-10-225-567A-233	Sequence 233, App
32	43	30.1	543	15	US-10-156-761-13485	Sequence 13485, A
33	43	30.1	571	9	US-09-815-242-11813	Sequence 11813, A
34	43	30.1	582	10	US-09-331-631A-22	Sequence 22, Appli
35	43	30.1	640	9	US-09-989-723-501	Sequence 501, App
36	43	30.1	640	9	US-09-989-723-501	Sequence 501, App
37	43	30.1	640	9	US-09-989-729-501	Sequence 501, App
38	43	30.1	640	9	US-09-989-727-501	Sequence 501, App
39	43	30.1	640	10	US-09-989-731-501	Sequence 501, App
40	43	30.1	640	10	US-09-989-732-501	Sequence 501, App
41	43	30.1	640	10	US-09-991-073-501	Sequence 501, App
42	43	30.1	640	10	US-09-909-320-292	Sequence 292, App
43	43	30.1	640	10	US-09-990-442-501	Sequence 501, App
44	43	30.1	640	10	US-09-991-163-501	Sequence 501, App
45	43	30.1	640	10	US-09-993-604-501	Sequence 501, App

ALIGNMENTS

RESULT 1
US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Oltersdorf, Tilman
; APPLICANT: Horne, William A.
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; FILE REFERENCE: 480140.42803
; CURRENT APPLICATION NUMBER: US/09/922.378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 94.4%; Score 135; DB 9; Length 204;
Best Local Similarity 96.2%; Pred. No. 5.5e-12;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRLFRMSDEFEGSKGL 27
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DB 140 NLWAAQRYGRLFRMTDEFEGSKGL 165

RESULT 2
US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US2002015631A1
; GENERAL INFORMATION:
; APPLICANT: Oltersdorf, Tilman
; APPLICANT: Horne, William A.
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE

; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-066-179-3

Query Match 94.4%; Score 135; DB 14; Length 204;
Best Local Similarity 96.2%; Pred. No. 5.5e-12;
Matches 25; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSKGL 27
|||||
Db 140 NLWAAQRYGRELRRMTDFEGSKGL 165

RESULT 3

US-10-059-261-258
; Sequence 258, Application US/10059261
; Publication No. US20030077826A1
; GENERAL INFORMATION:
; APPLICANT: EDELMAN, LENA
; APPLICANT: JACOTOT, ETIENNE DANIEL FRANCOIS
; APPLICANT: BRIAND, JEAN-PAUL
; TITLE OF INVENTION: CHIMERIC MOLECULES CONTAINING A MODULE ABLE TO TARGET
; TITLE OF INVENTION: SPECIFIC CELLS AND A MODULE REGULATING THE APOPTOTIC
; TITLE OF INVENTION: FUNCTION OF THE PERMEABILITY TRANSITION PORE COMPLEX
; TITLE OF INVENTION: (PTPC)
; FILE REFERENCE: 03495.0216
; CURRENT APPLICATION NUMBER: US/10/059,261
; CURRENT FILING DATE: 2002-08-29
; PRIOR APPLICATION NUMBER: 60/265,594
; PRIOR FILING DATE: 2001-02-02
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 258
; LENGTH: 25
; TYPE: PRT
; ORGANISM: Unknown Organism
; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: TOX peptide
US-10-059-261-258

Query Match 79.7%; Score 114; DB 15; Length 25;
Best Local Similarity 91.7%; Pred. No. 7.4e-10;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSKF 25
|||||
Db 1 NLWAAQRYGRELRRMSDFVDSFK 24

RESULT 4

US-09-922-378-2
; Sequence 2, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT

; ORGANISM: Homo sapiens
US-09-922-378-2

Query Match 79.7%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.1e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSKF 25
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Db 103 NLWAAQRYGRELRRMSDFVDSFK 126

RESULT 5

US-09-894-657-1
; Sequence 1, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: SF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 358673
; LIBRARY: SYNORAB01
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-894-657-1

Query Match 79.7%; Score 114; DB 9; Length 168;
Best Local Similarity 91.7%; Pred. No. 5.1e-09;
Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDFEGSKF 25
|||||
Db 103 NLWAAQRYGRELRRMSDFVDSFK 126

RESULT 6

US-09-894-657-7

FILE REFERENCE: ELITRA.011A
CURRENT APPLICATION NUMBER: US/09/815,242
CURRENT FILING DATE: 2001-03-21
PRIOR APPLICATION NUMBER: 60/191,078
PRIOR FILING DATE: 2000-03-21
PRIOR APPLICATION NUMBER: 60/206,848
PRIOR FILING DATE: 2000-05-23
PRIOR APPLICATION NUMBER: 60/207,727
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: 60/242,578
PRIOR FILING DATE: 2000-10-23
PRIOR APPLICATION NUMBER: 60/253,625
PRIOR FILING DATE: 2000-11-27
PRIOR APPLICATION NUMBER: 60/257,931
PRIOR FILING DATE: 2000-12-22
PRIOR APPLICATION NUMBER: 60/269,308
PRIOR FILING DATE: 2001-02-16
NUMBER OF SEQ ID NOS: 14110
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 5704
LENGTH: 426
TYPE: PRT
ORGANISM: Staphylococcus aureus
US-09-815-242-5704

Query Match 30.8%; Score 44; DB 9; Length 426;
Best Local Similarity 36.8%; Pred. No. 2.1e+02;
Matches 7; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

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core 46; DB 9; Length 946;
Pred. No. 2.4e+02;
5; Mismatches 13; Indels 0; Gaps 0;

Qy 3 ASDEFGSKGL 27
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Db LHVPDTFGHFDGV 237

RESULT 14
US-10-156-761-1151
Sequence 11541, Application US/10156761
Publication No. US20030119018A1
GENERAL INFORMATION:
APPLICANT: OMURA, SATOSHI
APPLICANT: IKEDA, HARUO
APPLICANT: ISHIKAWA, JUN
APPLICANT: HORIKAWA, HIROSHI
APPLICANT: SHIBA, TADAYOSHI
APPLICANT: SAKAKI, YOSHIYUKI
APPLICANT: HATTORI, MASAHIRA
TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
FILE REFERENCE: 249-262
CURRENT APPLICATION NUMBER: US/10/156,761
CURRENT FILING DATE: 2002-05-29
PRIOR APPLICATION NUMBER: JP 2001-204089
PRIOR FILING DATE: 2001-05-30
PRIOR APPLICATION NUMBER: JP 2001-272697
PRIOR FILING DATE: 2001-08-02
NUMBER OF SEQ ID NOS: 15109
SEQ ID NO 11541
LENGTH: 272
TYPE: PRT
ORGANISM: Streptomyces avermitilis
US-10-156-761-11541

Query Match 30.8%; Score 44; DB 15; Length 272;
Best Local Similarity 53.3%; Pred. No. 1.3e+02;
Matches 8; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

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Db 29 WIAAHGAELRRAD 43

RESULT 15
US-09-815-242-5704
Sequence 5704, Application US/09815242
Patent No. US20020061569A1
GENERAL INFORMATION:
APPLICANT: Haselbeck, Robert
APPLICANT: Ohlsen, Kari L.
APPLICANT: Zyskind, Judith W.
APPLICANT: Wall, Daniel
APPLICANT: Trawick, John D.
APPLICANT: Carr, Grant J.
APPLICANT: Yamamoto, Robert T.
APPLICANT: Xu, H. Howard
TITLE OF INVENTION: Identification of Essential Genes in
TITLE OF INVENTION: Prokaryotes

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OM protein - protein search, using sw model

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Title: US-09-544-664-56
Perfect score: 143
Sequence: 1 KNLWAAQRYGRELRRMSDEFGSKGL 27

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
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2: pir2: *
3: pir3: *
4: pir4: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	138	96.5	204	A55671	bad protein - mous
2	54	37.8	946	JC5575	inter-alpha-trypsi
3	53	37.1	223	D70760	hypothetical prote
4	53	37.1	946	S54354	inter-alpha-inhibi
5	52	36.4	370	S38185	2-dehydro-3-deoxy-
6	51	35.7	232	A42095	floral homeotic pr
7	50	35.0	374	C84338	spermidine/putresc
8	50	35.0	516	A86753	probable threonine
9	49	34.3	453	E83517	conserved hypothet
10	48.5	33.9	134	S40376	ig kappa chain - h
11	48.5	33.9	314	T02975	annexin P35 - maiz
12	48	33.6	206	T03635	transforming prote
13	48	33.6	220	F72289	oxido-reductase, so
14	48	33.6	526	T08545	threonine synthase
15	47.5	33.2	334	A39172	Antho-Ramide neur
16	47	32.9	597	F82308	oxalacetate decar
17	47	32.9	967	F82668	oxoglutarate dehyd
18	47	32.9	5138	B96695	hypothetical prote
19	46.5	32.5	314	T02961	annexin P33 - maiz
20	46.5	32.5	435	A44308	Antho-Ramide prec
21	46.5	32.5	1140	T09486	hypothetical prote
22	46	32.2	165	S59899	chlorocruorin chal
23	46	32.2	399	T35440	probable polyamine
24	46	32.2	946	IVHU2	inter-alpha-trypsi
25	46	32.2	1164	T24806	hypothetical prote
26	46	32.2	1378	A81393	DNA-directed RNA p
27	45.5	31.8	261	G69510	conserved hypothet
28	45.5	31.8	287	S43852	neuropeptide Pol-R
29	45.5	31.8	327	AF2859	conserved hypothet

30 45.5 31.8 327 2 D97636 probable secreted
31 45.5 31.8 562 2 C71473 hypothetical prote
32 45.5 31.8 905 2 G83314 NADH dehydrogenase
33 45 31.5 273 2 S06736 photosystem II oxy
34 45 31.5 273 2 A32287 manganese-stabilizi
35 45 31.5 295 2 F83201 conserved hypothet
36 45 31.5 346 2 H95406 conserved hypothet
37 45 31.5 591 2 B44465 sodium ion pump ox
38 45 31.5 591 2 A80509 oxalacetate decar
39 45 31.5 591 2 A80909 oxalacetate decar
40 45 31.5 596 2 A28088 oxalacetate decar
41 45 31.5 715 2 S52675 probable membrane
42 45 31.5 864 1 VCLJG4 env polypeptide -
43 45 31.5 915 2 B59433 chromosome 5 GAP-1
44 45 31.5 1199 2 T23005 hypothetical prote
45 45 31.5 1217 2 T22672 hypothetical prote

ALIGNMENTS

RESULT 1

A55671
bad protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999
C:Accession: A55671
R:Yang, E.; Zha, J.; Jockel, J.; Boise, L.H.; Thompson, C.B.; Korsmeyer, S.J.
Cell 80, 285-291, 1995
A:Title: Bad, a heterodimeric partner for Bcl-x-L and Bcl-2, displaces Bax and promot
A:Reference number: A55671; MUID:95136361; PMID:7834748
A:Accession: A55671
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-204 <NAK>
C:Cross-references: GB:I37296; NID:g6397778; PIDN:AAA64465.1; PID:g6397779
C:Keywords: heterodimer

Query Match 96.5%; Score 138; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. NO. 9.3e-13;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRRMSDEFGSKGL 27
|||||
DB 140 NLWAAQRYGRELRRMSDEFGSKGL 165

RESULT 2

JC5575
inter-alpha-trypsin inhibitor heavy chain 2 - golden hamster
C:Species: Mesocricetus auratus (golden hamster)
C:Date: 23-Sep-1997 #sequence_revision 23-Sep-1997 #text_change 20-Jun-2000
C:Accession: JC5575; PC4485
R:Nakatani, T.; Suzuki, Y.; Yamamoto, T.; Sinohara, H.
J. Biochem. 122, 71-82, 1997
A:Title: Molecule cloning and sequencing of cDNAs encoding three heavy-chain precurs
sin inhibitor heavy chain family.
A:Reference number: JC5574; MUID:97420688; PMID:9276673
A:Accession: JC5575
A:Molecule type: mRNA
A:Residues: 1-946 <NAK>
A:Cross-references: DDBJ:D89286; NID:g1694689; PIDN:BAAL1939.1; PID:g1694690
A:Experimental source: liver
A:Accession: PC4485
A:Molecule type: protein
A:Residues: 55-64;140-146;151-156;424-447;500-528;577-605 <NA2>
C:Comment: In the plasma three inter-alpha-trypsin inhibitor heavy chains 1, 2 and 3
that the complexes play important role for pancreatic cancer.
C:Superfamily: inter-alpha-trypsin inhibitor complex component II
F;261-264,717-916/Disulfide bonds: #status predicted

Query Match 37.8%; Score 54; DB 2; Length 946;
Best Local Similarity 34.6%; Pred. No. 9.9;

A;Title: The complete sequence of a 6794 bp segment located on the right arm of chromosome 10p11.23, containing the human APOA2 gene and 5' flanking region.
A;Reference number: S3185; MUID:94078675; PMID:8256522
A;Accession: S3185
A;Status: translation not shown
A:Molecule type: DNA
A;Residues: 1-370 <DOI>
A;Cross-references: GB:120296; NID:g311101; PIDN:AAA56507.1; PID:g311102
R:Aljinovic, G.; Pohl, F.M.; Pohl, T.M.
submitted to the Protein Sequence Database, August 1994
A;Reference number: S45906
A;Accession: S46126
A:Molecule type: DNA
A;Residues: 1-370 <AL>
A;Cross-references: EMBL:336118; NID:g536664; PIDN:CAA85212.1; PID:g536665; MIPS:YBR2
R:Algie, M.; Baciuet, M.C.; Barthe, C.; Biteau, N.; Crouzet, M.; Doignon, F.
submitted to the Protein Sequence Database, August 1994
A;Reference number: S45940
A;Accession: S46130
A:Molecule type: DNA
A;Residues: 1-370 <ATG>
A;Cross-references: EMBL:336118; NID:g536664; PIDN:CAA85212.1; PID:g536665; MIPS:YBR2
R:Kuenzler, M.; Paravicini, G.; Egli, C.M.; Irniger, S.; Braus, G.H.
Gene 113, 67-74, 1992
A;Title: Cloning, primary structure and regulation of the ARO4 gene, encoding the tyrosine aminotransferase
A;Reference number: JN0322; MUID:92225349; PMID:1348717
A;Accession: JN0322
A:Molecule type: DNA
A;Residues: 1-204,208-370 <KUE>
A;Cross-references: EMBL:X61107
R:Kuenzler, M.; Balmeili, T.; Egli, C.M.; Paravicini, G.; Braus, G.H.
J. Bacteriol. 175, 5548-5558, 1993
A;Title: Cloning, primary structure, and regulation of the HIS7 gene encoding a bifunctional histidinol-phosphatase
A;Reference number: A48651; MUID:93374850; PMID:8366040
A;Accession: B48651
A;Status: preliminary
A:Molecule type: DNA
A;Residues: 352-370 <KU2>
A;Cross-references: GB:X61107
C;Comment: This enzyme catalyzes the condensation of phosphoenolpyruvate and D-erythrulose to form 4-keto-7-phosphoheptulon.
C;Genetics:
A;Gene: SGD:ARO4
A;Cross-references: SGD:S0000453; MIPS:YBR249C
A;Map position: 2R
C;Function:
A;Description: aldehyde-lyase; carbon-carbon lyase
A;Pathway: aromatic amino acid biosynthesis; shikimate pathway
A;Note: first step in shikimate pathway
C;Superfamily: phospho-2-dehydro-3-deoxyheptonate aldolase
C;Keywords: aldehyde-lyase, aromatic amino acid biosynthesis; carbon-carbon lyase; cytochrome b5
Query Match 36.4%; Score 52; DB 2: Length 370;
Best Local Similarity 47.6%; Pred. No. 7.5;
Matches 10; Conservative 5; Mismatches 6; Indels 0; Gaps 0;
QY 2 NLAAQRYGRELRLMSDEEG 22
:|||||:||||:
Db 80 DLEAAQRYALRLKLSDLEK 100
RESULT 6
A42095
N;Alternate names: homeotic protein APTTALA3 (AP3) - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 04-Mar-1993 #sequence_revision 18-Nov-1994 #text_change 21-Jul-2000
C;Accession: A42095; S52633; T47593
R:Jack, T.; Brockman, L.L.; Meyerowitz, E.M.
Cell 68, 683-697, 1992
A;Title: The homeotic gene APTTALA3 of Arabidopsis thaliana encodes a MADS box and is involved in floral homeotic protein APTTALA3 (AP3) - Arabidopsis thaliana
A;Reference number: A42095; MUID:92154682; PMID:1346756
A;Accession: A42095
A;Status: preliminary
A:Molecule type: mRNA

C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001

C:Accession: A96753
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alon
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar,
ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000

A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim,
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Mafti, R.; Marzia
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallo
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.

A>Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21036719; PMID:11130712
A:Accession: A96753
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-516 <STO>
A:Cross-references: GB:AE005173; NID:g5903070; PIDN:AAD55628.1; GSPDB:GN00141
C:Genetics:
A:Gene: F3N23.1
A:Map position: 1

Query Match 35.0%; Score 50; DB 2; Length 516;
Best Local Similarity 35.3%; Pred. No. 21;
Matches 12; Conservative 7; Mismatches 7; Indels 8; Gaps 1;

QY 2 NLWAAQRYGRELRLMSD-----EFEGSRKGL 27
|||::||::||::||::||
DB 163 NLFWAEFRKGQLNMNLWKHCGISHTGSFKDL 196

RESULT 9
E83517
C:Species: Pseudomonas aeruginosa
C: conserved hypothetical protein PA1031 [imported] - Pseudomonas aeruginosa (strain PAO
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
R:Accession: E83517
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.;
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; L
Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000

A:Title: Complete genome sequence of Pseudomonas aeruginosa PAO1, an opportunistic pa
A:Reference number: A82950; MUID:20437337; PMID:10984043
A:Accession: E83517
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-453 <STO>
A:Cross-references: GB:AE004535; GB:AE004091; NID:g9946936; PIDN:AAG04420.1; GSPDB:GN
A:Experimental source: strain PAO1
C:Genetics:
A:Gene: PAI031

Query Match 34.3%; Score 49; DB 2; Length 453;
Best Local Similarity 55.6%; Pred. No. 25;
Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;

QY 4 WAAQRYGR--ELRRMSDE 19
||||| || | |||| :: |
DB 65 WASTRGREELRLASE 82

RESULT 10
S40376
I9 kappa chain - human
C:Species: Homo sapiens (man)
C:Date: 06-Mar-1994 #sequence_revision 26-May-1995 #text_change 21-Jan-2000
R:Accession: S40376
R:Klein, R.; Jaenichen, R.; Zachau, H.G.
Eur. J. Immunol. 23, 3248-3271, 1993

A:Title: Expressed human immunoglobulin chi genes and their hypermutation.
A:Reference number: S40312; MUID:94080891; PMID:8258341
A:Accession: S40376
A>Status: preliminary; translation not shown

Db 169 RETRANKEQGRSGK 184

RESULT 13
F72289
oxidoreductase, sol/devB family - Thermotoga maritima (strain MSB8)
C:Species:Thermotoga maritima
C>Date:11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 21-Jul-2000
C:Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwinn, M.L.; Dodson, R.J.; Haft, D.H.; Hic,
Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson,
C.M.
Nature 399, 323-329, 1999
A:title:Evidence for lateral gene transfer between Archaea and Bacteria from genome
A:Reference number: A72200; MOID:99287316; PMID:10360571
A:Accession: F72289
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-220 <ARN>
A:Cross-references: GB:AE001772; GB:AE000512; NID:g4981693; PIDN:AAD36230.1; PID:g4981693
A:Experimental source: strain MSB8
C:Genetics:
A:Gene: TWI154
C:Superfamily: yeast SOL3 protein

Query Match 33.6%; Score 48; DB 2; Length 220;
Best Local Similarity 34.8%; Pred. No. 17;
Matches 8; Conservative 8; Mismatches 7; Indels 0; Gaps 0;

QY 5 AAQRYGRELRRMSDEFGSFGL 27
|::|||::|::|::|
Db 111 ACEKYERISATDQDFLAILGM 133

RESULT 14
T08545
threonine synthase (EC 4.2.3.1) precursor - Arabidopsis thaliana
N:Alternate names: protein F27B13.80
C:Species:Arabidopsis thaliana (mouse-ear cress)
C>Date:11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 03-Jun-2002
C:Accession: T08545; S71362; S74307
R:Bevan, M.; Zimmermann, W.; Gruenisen, A.; Wambutt, R.; Bancroft, I.; Meyes, H.W.;
submitted to the Protein Sequence Database, May 1999
A:Reference number: Z16442
A:Accession: T08545
A:Molecule type: DNA
A:Residues: 1-526 <BEV>
A:Cross-references: EMBL:AL050352; GSPDB:GN00062; ATSP:F27B13.80
A:Experimental source: cultivar Columbia; BAC clone F27B13
P:Curien, G.; Dumas, R.; Ravanel, S.; Douce, R.
FEBS Lett. 390, 85-90, 1996
A:title:Characterization of an Arabidopsis thaliana cDNA encoding an S-adenosylmethil
A:Reference number: S71362; MOID:96314555; PMID:8706836
A:Accession: S71362
A:Molecule type: mRNA
A:Residues: 1/1, 3-526 <CUR>
A:Cross-references: EMBL:L41666; NID:g1448916; PIDN:AAB04607.1; PID:g1448917
A:Accession: S74307
A:Molecule type: protein
A:Residues: 40-54 <CUI>
C:Genetics:
A:Gene: ATSP:F27B13.80
A:Map position: 4
A:Genome: nuclear
C:Keywords: carbon-oxygen lyase; chloroplast
F:1-39/Domain: transit peptide (chloroplast) #status predicted <TNP>
F:40-526/Product: threonine synthase #status experimental <MAT>

Query Match 33.6%; Score 48; DB 2; Length 526;
Best Local Similarity 35.3%; Pred. No. 42;
Matches 12; Conservative 6; Mismatches 8; Indels 8; Gaps 1;

QY 2 NLWAAQRYGRELRRMSD-----EFGSFKGL 27

Db 172 NLFWAERFGKQFLGMNDLWVKHCGISHTSFKDL 205

RESULT 15

A39172
 Antho-RFamide neuropeptide 19 repeat precursor - sea anemone (*Calliactis parasitica*)
 C:Species: *Calliactis parasitica*
 C:Date: 07-Feb-1992 #sequence_revision 07-Feb-1992 #text_change 21-Jul-2000
 C:Accession: A39172
 E:Barner, D.; Schmutzler, C.; Diekhoff, D.; Grimmelikhuijzen, C.J.P.
 Proc. Natl. Acad. Sci. U.S.A. 88, 2553-2559, 1991
 A:Title: Primary structure of the precursor for the sea anemone neuropeptide Antho-RFamide
 A:Reference number: A39172; MUID:91172845; PMID:1706527
 A:Accession: A39172
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-334 <DAR>
 A:Cross-references: GB:MS9166; NID:gl56133; PID:AAA27878.1; PID:gl56134
 C:Keywords: neuropeptide

```
Query Match      33.2%; Score 47.5; DB 2; Length 334;
Best Local Similarity 44.0%; Pred. No. 31;
Matches 11; Conservative 3; Mismatches 10; Indels 1; Gaps 1;
```

QY 1 KNLWAAQRYGRELR-RMSDEFFEGSF 24

89 KRRYVPGRYGRFQGRFGREFQGRF 113

Search completed: September 15, 2003, 17:27:04
Job time : 13.15 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:55 : Search time 6.36429 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-56

Perfect score: 143

Sequence: 1 KNLWAAQRYGRELRRMSDFEGSKGL 27

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	138	96.5	204	1 BAD_MOUSE	Q61337 mus musculus
2	138	96.5	205	1 BAD_RAT	Q35147 rattus norv
3	114	79.7	168	1 ITH2_HUMAN	Q92934 homo sapien
4	54	37.8	946	1 ITH2_MESAU	P97279 mesocricetu
5	53	37.1	946	1 ITH2_MOUSE	Q61703 mus musculus
6	52.5	36.7	506	1 MATK_LEDPA	Q62992 ledum palus
7	52.5	36.7	506	1 MATK_RHOFR	Q62984 rhododendro
8	52.5	36.7	506	1 MATK_RHOTS	Q62991 rhododendro
9	52	36.4	370	1 AROG_YEAST	P32449 saccharomyc
10	51	35.7	232	1 AP3_ARATH	P35632 arabidopsis
11	51	35.7	851	1 CE05_MOUSE	Q8K2H3 mus musculus
12	49	34.3	453	1 RMUC_PSEAE	Q914U3 pseudomonas
13	48	33.6	205	1 RAS3_RHIRA	P22280 rhizomucor
14	48	33.6	220	1 6PGL_THEMA	Q9X0N8 thermotoga
15	48	33.6	519	1 THRC_SOLUTU	Q9MT28 solanum tub
16	48	33.6	526	1 THRC_ARATH	Q9S7B5 arabidopsis
17	47.5	33.2	334	1 FWRA_CALPA	Q01133 callitactis
18	47.5	33.2	507	1 MATK_LOIPR	Q47169 loiseleuria
19	47	32.9	198	1 BIM_HUMAN	Q43521 homo sapien
20	46.5	32.5	429	1 FMR2_ANTEL	Q16994 antiopeura
21	46.5	32.5	435	1 FMR1_ANTEL	P10419 antiopeura
22	46	32.2	946	1 ITH2_HUMAN	P19823 homo sapien
23	46	32.2	1378	1 RPOB_CAMJE	Q46124 campylobact
24	45.5	31.8	287	1 PRFA_POLPE	P21259 polyorchis
25	45	31.5	273	1 PSBO_ANASP	P13907 anabaena sp
26	45	31.5	328	1 SNF4_KLULA	Q9P869 kluyveromyc
27	45	31.5	590	1 DCOA_SALTY	Q03030 salmonella
28	45	31.5	595	1 DCOA_KLEPN	P13187 klebsiella
29	45	31.5	653	1 HT2A_HUMAN	Q13049 homo sapien
30	45	31.5	865	1 ENV_SIVAT	P05886 simian immu
31	45	31.5	915	1 CE05_HUMAN	Q9NYF5 homo sapien
32	45	31.5	1535	1 LML1_CAEEL	Q18823 caenorhabdi
33	44.5	31.1	506	1 MATK_GAUPR	Q95GJ0 gaultheria

34	44.5	31.1	512	1 MATK_LILTS	O9GIG3 lilium tsain
35	44.5	31.1	907	1 NUOC_ECOLI	P33602 escherichia
36	44.5	31.1	907	1 NUOC_SALTY	P33900 salmonella
37	44	30.8	196	1 BIM_MOUSE	O54918 mus musculus
38	44	30.8	196	1 BIM_RAT	O88498 rattus norv
39	44	30.8	262	1 END8_ECO57	O8XG66 escherichia
40	44	30.8	262	1 END8_ECOLI	P50465 escherichia
41	44	30.8	262	1 END8_SALTY	O8Z8D2 salmonella
42	44	30.8	262	1 END8_SALTY	Q8ZQ66 salmonella
43	44	30.8	629	1 SYM_THEMEA	O33925 thermotoga
44	44	30.8	768	1 ENV_SIVAI	P27757 simian immu
45	44	30.8	877	1 ENV_SIVAG	P27977 simian immu

ALIGNMENTS

RESULT 1	BAD_MOUSE	STANDARD;	PRT;	204 AA.
ID	Q61337;			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	28-FEB-2003 (Rel. 41, Last annotation update)			
DE	Bcl-2-antagonist of cell death (BAD) (Bcl-2 binding component			
DE	6) (Bcl-xL/Bcl-2 associated death promoter).			
GN	BAD OR BCL6.			
OS	Mus musculus (Mouse).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.			
OX	NCBI_TaxID=10090;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Brain, and Thymus;			
RC	MEDLINE=95136361; PubMed=7834748;			
RA	Yang E., Zhu J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;			
RT	"Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and			
RT	promotes cell death."			
RL	Cell 80:285-291(1995).			
RN	[2]			
RP	PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.			
RC	MEDLINE=98022383; PubMed=9381178;			
RA	Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;			
RT	"Interleukin-3-induced phosphorylation of BAD through the protein			
RT	kinase Akt."			
RL	Science 278:687-689(1997).			
RN	[3]			
RP	MUTAGENESIS OF SERINE RESIDUES.			
RC	MEDLINE=20403302; PubMed=10949026;			
RA	Datta S.R., Katsov A., Hu L., Petros A., Fesik S.W., Yaffe M.B.,			
RT	Greenberg M.E.;			
RT	"14-3-3 proteins and survival kinases cooperate to inactivate BAD by			
RT	BH3 domain phosphorylation."			
RL	Mol. Cell 6:41-51(2000).			
CC	CC -!- FUNCTION: Promotes cell death. Successfully competes for the			
CC	binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level			
CC	of heterodimerization of these proteins with BAX. Can reverse the			
CC	death repressor activity of Bcl-x(L), but not that of Bcl-2.			
CC	Appears to act as a link between growth factor receptor signaling			
CC	and the apoptotic pathways.			
CC	CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-			
CC	x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).			
CC	The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.			
CC	CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon			
CC	phosphorylation, locates to the cytoplasm.			
CC	CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND			
CC	BAX for their pro-apoptotic activity and for their interaction			
CC	with anti-apoptotic members of the Bcl-2 family.			
CC	CC -!- PTM: Phosphorylated on Ser-112 in response to survival stimuli.			
CC	Subsequent phosphorylation on Ser-136 promotes heterodimerization			
CC	with 14-3-3 proteins. This interaction then facilitates the			
CC	phosphorylation at Ser-155, a site within the BH3 domain, leading			
CC	to the release of Bcl-x(L) and the promotion of cell survival.			

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CC Ser-136 is the major site of AKT/PKB phosphorylation, Ser-155 the
CC major site of protein kinase A (CAPK) phosphorylation.
CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; L37296; AAA64465.1; -
CC PIR; A55671; A55671.
CC HSSP; Q92934; IG5J.
CC MGD; MGI:1096330; Bad.
CC InterPro; IPR000712; Bcl2_BH.
CC PROSITE; PS01259; BH3; FALSE_NEG.
CC Apoptosis; Phosphorylation.
CC DOMAIN 147 161 BH3.
CC MOD_RES 112 112 PHOSPHORYLATION (BY PKA AND PKB).
CC MOD_RES 136 136 PHOSPHORYLATION (BY PKA AND PKB).
CC MOD_RES 155 155 PHOSPHORYLATION (BY PKA AND PKB).
CC MUTAGEN 112 112 S->A: NO PHOSPHORYLATION.
CC MUTAGEN 136 136 S->A: NO PHOSPHORYLATION.
CC MUTAGEN 155 155 S->A: NO PHOSPHORYLATION; INTERACTS WITH
CC BCL-X(L).
CC SEQUENCE 204 AA; 22080 MW; 6C2BA910205053F7 CRC64;
CC -----
CC Query Match 96.5%; Score 138; DB 1; Length 204;
CC Best Local Similarity 100.0%; Pred. No. 1.8e-13;
CC Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CC -----
CC QY 2 NLWAAQRYGRELRLMSDFGSGPKGL 27
CC 140 NLWAAQRYGRELRLMSDFGSGPKGL 165
CC -----
CC RESULT 2
CC BAD_RAT
CC ID BAD_RAT STANDARD; PRT; 205 AA.
CC AC O35147; O70256; Q9JHX1;
CC DT 16-OCT-2001 (Rel. 40, Created)
CC DT 16-OCT-2001 (Rel. 40, Last sequence update)
CC DT 28-FEB-2003 (Rel. 41, Last annotation update)
CC DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component
CC 6) (Bcl-xL/Bcl-2 associated death promoter).
CC GN BAD.
CC OS Rattus norvegicus (Rat).
CC OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
CC OX NCBI_TaxID=10116;
CC RN [1]
CC RP SEQUENCE FROM N.A., AND MUTAGENESIS OF SER-113 AND SER-137.
CC RC TISSUE-Ovary;
CC RA MEDLINE=98034386; PubMed=9369453;
CC RA Hsu S.Y., Kaipia A., Zhu L., Hsueh A.J.W.;
CC RT "Interference of BAD (Bcl-xL/Bcl-2-associated death promoter)-induced
CC RT apoptosis in mammalian cells by 14-3-3 isoforms and P11.";
CC RL Mol. Endocrinol. 11:1858-1867(1997).
CC RN [2]
CC RP SEQUENCE FROM N.A.
CC RC TISSUE=Brain;
CC RX MEDLINE=98194755; PubMed=9535132;
CC RA D'Agata V., Magro G., Travali S., Musco S., Cavallaro S.;
CC RT "Cloning and expression of the programmed cell death regulator BAD in
CC RT the rat brain.";
CC RL Neurosci. Lett. 243:137-140(1998).
CC RN [3]
CC RP SEQUENCE FROM N.A. (ISOFORMS ALPHA AND BETA).
CC RC TISSUE=Brain;
CC RX MEDLINE=21109372; PubMed=11161472;

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RA Hanner S., Arumae U., Yu L.-Y., Sun Y.-P., Saarma M., Lindholm D.;
RA "Functional characterization of two splice variants of rat BAD and
RA their interaction with Bcl-w in sympathetic neurons.";
RA Mol. Cell. Neurosci. 17:97-106(2001).
CC -!- FUNCTION: Promotes cell death. Successfully competes for the
CC binding to Bcl-x(L). Bcl-2 and Bcl-w thereby affecting the level
CC of heterodimerization of these proteins with BAX. Can reverse the
CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
CC similarity). Appears to act as a link between growth factor
CC receptor signaling and the apoptotic pathways.
CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10. The Ser-
CC 113/Ser-137 phosphorylated form binds 14-3-3 proteins.
CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
CC phosphorylation, locates to the cytoplasm (By similarity).
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=Alpha;
CC IsoId=O35147-1; Sequence=Displayed;
CC Name=Beta;
CC IsoId=O35147-2; Sequence=VSP_000534;
CC -!- TISSUE SPECIFICITY: Expressed in all tissues tested, including
CC brain, liver, spleen and heart. In the brain, restricted to
CC epithelial cells of the choroid plexus. Isoform alpha is the more
CC abundant form.
CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
CC BAX for their pro-apoptotic activity and for their interaction
CC with anti-apoptotic members of the Bcl-2 family.
CC -!- PTM: Phosphorylated on Ser-113 in response to survival stimuli.
CC Subsequent phosphorylation on Ser-137 promotes heterodimerization
CC with 14-3-3 proteins. This interaction then facilitates the
CC phosphorylation at Ser-156, a site within the BH3 domain, leading
CC to the release of Bcl-x(L) and the promotion of cell survival.
CC Ser-137 is the major site of AKT/PKB phosphorylation, Ser-156 the
CC major site of protein kinase A (CAPK) phosphorylation (By
CC similarity).
CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF003523; AAC53374.1; -
CC EMBL; AF031227; AAC15100.1; -
CC EMBL; AF279910; AAF91427.1; -
CC EMBL; AF279911; AAF91428.1; -
CC HSSP; Q92934; IG5J.
CC InterPro; IPR000712; Bcl2_BH.
CC PROSITE; PS01259; BH3; FALSE_NEG.
CC Apoptosis; Phosphorylation; Alternative splicing.
CC DOMAIN 148 162 BH3.
CC MOD_RES 113 113 PHOSPHORYLATION (BY PKA AND PKB)
CC (BY SIMILARITY).
CC MOD_RES 137 137 PHOSPHORYLATION (BY PKA AND PKB)
CC (BY SIMILARITY).
CC MOD_RES 156 156 PHOSPHORYLATION (BY PKA AND PKB)
CC (BY SIMILARITY).
CC VARSPLOC 166 205 LPRPKSAGTACMRCOSASWTRIIQSWDRNLKGGSTPSQ
CC -> EELTYSVEFLPVRAIMEGAPLWSQSFPHLPPTPP
CC /FTID=VSP_000534.
CC MUTAGEN 113 113 S->A: NO EFFECT ON HETERODIMERIZATION
CC WITH 14-3-3 PROTEINS.
CC MUTAGEN 137 137 S->A: NO HETERODIMERIZATION WITH 14-3-3
CC PROTEINS. NO EFFECT ON HETERODIMERIZATION
CC WITH BCL2 NOR WITH PROTEIN P11.
CC CONFLICT 29 34 SDAGGR -> ERRGRK (IN REF. 1).
CC SEQUENCE 205 AA; 22228 MW; 7AFA71DAE9CF4A81 CRC64;

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Best Local Similarity 91.7%; Pred. No. 5.8e-10; Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFEFSFK 25
 Db 103 NLWAAQRYGRELRLMSDEFEFSFK 126

RESULT 4
 ID ITH2_MESAU STANDARD; PRT; 946 AA.
 AC P97279;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DE Inter-alpha-trypsin inhibitor heavy chain H2 precursor (ITI heavy chain H2) (Inter-alpha-inhibitor heavy chain 2) (HC2).
 GN ITIH2.
 OS Mesocricetus auratus (Golden hamster).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae; Mesocricetus.
 OX NCBI_TaxID=10036;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX MEDLINE=97420688; PubMed=9276673;
 RA Nakatani T., Suzuki Y., Yamamoto T., Sinohara H.;
 RT "Molecular cloning and sequencing of cDNAs encoding three heavy-chain precursors of the inter-alpha-trypsin inhibitor in Syrian hamster: implications for the evolution of the inter-alpha-trypsin inhibitor heavy chain family";
 RT J. Biochem. 122:71-82(1997).
 RL [2]
 RN SEQUENCE OF 55-64; 140-146; 151-156; 424-447; 500-528 AND 577-605, AND SUBUNITS.
 RP TISSUE=Plasma;
 RX MEDLINE=97018241; PubMed=8864857;
 RA Yamamoto T., Yamamoto K., Sinohara H.;
 RT "Inter-alpha-trypsin inhibitor and its related proteins in Syrian hamster urine and plasma";
 RL J. Biochem. 120:145-152(1996).
 CC -I- FUNCTION: MAY ACT AS A CARRIER OF HYALURONAN IN SERUM OR AS A BINDING PROTEIN BETWEEN HYALURONAN AND OTHER MATRIX PROTEIN, INCLUDING THOSE ON CELL SURFACES IN TISSUES TO REGULATE THE LOCALIZATION, SYNTHESIS AND DEGRADATION OF HYALURONAN WHICH ARE ESSENTIAL TO CELLS UNDERGOING BIOLOGICAL PROCESSES (BY SIMILARITY).
 CC -I- SUBUNIT: I-ALPHA-1 PLASMA PROTEASE INHIBITORS ARE ASSEMBLED FROM ONE OR TWO HEAVY CHAINS (H1, H2 OR H3) AND ONE LIGHT CHAIN, BIKUNIN. INTER-ALPHA-INHIBITOR (I-ALPHA-I) IS COMPOSED OF H1, H2 AND BIKUNIN, INTER-ALPHA-LIKE INHIBITOR (I-ALPHA-LI) OF H2 AND BIKUNIN, AND PRE-ALPHA-INHIBITOR (P-ALPHA-I) OF H3 AND BIKUNIN.
 CC -I- PTM: HEAVY CHAINS ARE INTERLINKED WITH BIKUNIN VIA A CHONDROITIN 4-SULFATE BRIDGE TO THE THEIR C-TERMINAL ASPARTATE (BY SIMILARITY).
 CC -I- SIMILARITY: BELONGS TO THE ITIH FAMILY.
 CC -I- SIMILARITY: Contains 1 WFPA domain.
 CC -----
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 CC -----
 CC EMBL; D89286; BAA13939.1; -
 CC PIR; JC5575; JC5575
 CC InterPro; IPR006587; VIT.
 CC InterPro; IPR002035; VWF_A.
 CC Pfam; PF00092; vwa; 1.
 CC SMART; SM00609; VIT; 1.

DR SMART; SM00327; VWA; 1.
 DR PROSITE; PS0234; VWF_A; 1.
 KW Serine protease inhibitor; Repeat; Signal; Multigene family;
 KW Glycoprotein.
 FT SIGNAL 1 18 POTENTIAL.
 FT PROPEP 19 54 BY SIMILARITY.
 FT CHAIN 55 702 INTER-ALPHA-TRYPSIN INHIBITOR HEAVY CHAIN H2.
 FT PROPEP 703 946 BY SIMILARITY.
 FT DOMAIN 308 468 VWF_A.
 FT CARBOHYD 118 118 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 263 263 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 445 445 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 578 578 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT BINDING 702 702 CHONDROITIN 4-SULFATE, CROSS-LINK SITE (BY SIMILARITY).
 FT CONFLICT 510 510 V -> Y (IN REF. 2).
 FT CONFLICT 595 595 E -> I (IN REF. 2).
 SQ SEQUENCE 946 AA; 106580 MW; CA8BF565458E7B2E CRC64;

Query Match 37.8%; Score 54; DB 1; Length 946;
 Best Local Similarity 34.6%; Pred. No. 3.7;
 Matches 9; Conservative 5; Mismatches 12; Indels 0; Gaps 0;

QY 2 NLWAAQRYGRELRLMSDEFEFSFKGL 27
 Db 212 NWVLEQGMFLHVPDTFEGHFGV 237

RESULT 5
 ID ITH2_MOUSE STANDARD; PRT; 946 AA.
 AC Q61703;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DE Inter-alpha-trypsin inhibitor heavy chain H2 precursor (ITI heavy chain H2) (Inter-alpha-inhibitor heavy chain 2).
 DE ITIH2.
 GN Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6N; TISSUE=Liver;
 RX MEDLINE=95194326; PubMed=7534067;
 RA Chan P., Rislter J.-L., Raguenez G., Salier J.-P.;
 RT "The three heavy-chain precursors for the inter-alpha-inhibitor family in mouse: new members of the multicopper oxidase protein group with differential transcription in liver and brain";
 RL Biochem. J. 306:505-512(1995).
 CC -I- FUNCTION: MAY ACT AS A CARRIER OF HYALURONAN IN SERUM OR AS A BINDING PROTEIN BETWEEN HYALURONAN AND OTHER MATRIX PROTEIN, INCLUDING THOSE ON CELL SURFACES IN TISSUES TO REGULATE THE LOCALIZATION, SYNTHESIS AND DEGRADATION OF HYALURONAN WHICH ARE ESSENTIAL TO CELLS UNDERGOING BIOLOGICAL PROCESSES (BY SIMILARITY).
 CC -I- SUBUNIT: I-ALPHA-1 PLASMA PROTEASE INHIBITORS ARE ASSEMBLED FROM ONE OR TWO HEAVY CHAINS (H1, H2 OR H3) AND ONE LIGHT CHAIN, BIKUNIN. INTER-ALPHA-INHIBITOR (I-ALPHA-I) IS COMPOSED OF H1, H2 AND BIKUNIN, INTER-ALPHA-LIKE INHIBITOR (I-ALPHA-LI) OF H2 AND BIKUNIN, AND PRE-ALPHA-INHIBITOR (P-ALPHA-I) OF H3 AND BIKUNIN.
 CC -I- TISSUE SPECIFICITY: EXPRESSED IN BOTH LIVER AND BRAIN.
 CC -I- PTM: HEAVY CHAINS ARE INTERLINKED WITH BIKUNIN VIA A CHONDROITIN 4-SULFATE BRIDGE TO THE THEIR C-TERMINAL ASPARTATE (BY SIMILARITY).
 CC -I- SIMILARITY: BELONGS TO THE ITIH FAMILY.
 CC -I- SIMILARITY: Contains 1 VWF_A domain.
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OX eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 RN NCBI_taxid=3702;
 RP [1]
 RC SEQUENCE FROM N.A.
 RX TISSUE-Petal;
 RA Jack T., Brockman L.L., Meyerowitz E.M.;
 RT "The homeotic gene APETALA3 of Arabidopsis thaliana encodes a MADS
 box and is expressed in petals and stamens.";
 RL Cell 68:683-697(1992).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Landsberg erecta;
 RX MEDLINE=95036018; PubMed=7948993;
 RA Okamoto H., Yano A., Shiraishi H., Okada K., Shimura Y.;
 RT "Genetic complementation of a floral homeotic mutation, apetal3,
 with an Arabidopsis thaliana gene homologous to DEFICIENS of
 Antirrhinum majus";
 RL Plant Mol. Biol. 26:465-472(1994).
 RN [3]
 RP SEQUENCE FROM N.A., AND VARIANTS.
 RC STRAIN=cv. Bla-1, cv. Bretagny, cv. Bs-1, cv. Bu-0, cv. Bu-2,
 cv. Chi-1, cv. Co-1, cv. Columbia, cv. Corsacalla-1, cv. Cvi-0,
 cv. Gr-3, cv. J1-1, cv. Kas-1, cv. Kent, cv. Landsberg erecta,
 cv. Li-3, cv. Li-8, and cv. Lisse;
 RX MEDLINE=99126449; PubMed=9927474;
 RA Purganan M.D., Sudduth J.I.;
 RT "Molecular population genetics of floral homeotic loci: departures
 from the equilibrium-neutral model at the APETALA3 and PISTILLATA
 genes of Arabidopsis thaliana.";
 RL Genetics 151:839-848(1999).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=21016720; PubMed=11130713;
 RA Salanoubat M., Lencke K., Rieger M., Ansoerge W., Unseld M.,
 Farmann B., Valle G., Bloeker H., Perez-Alonso M., Obermaier B.,
 Dessey M., Boutry M., Grivell L.A., Mache R., Puigdomenech P.,
 De Simone V., Choisine N., Artiguenave F., Robert C., Brottier P.,
 Wincker P., Catolico L., Weissenbach J., Saurin W., Quetier F.,
 Schaefer M., Mueller-Auer S., Gabel C., Fuchs M., Benes V.,
 Wurmbach E., Drzonek H., Erfle H., Jordan N., Bangert S.,
 Wiedemann R., Kranz H., Voss H., Holland R., Brandt P., Nvakatura G.,
 Verzi A., D'Angelo M., Pallavicini A., Toppo S., Simonati B.,
 Conrad A., Hornischer K., Kauer G., Loehner T.-H., Nordiek G.,
 Reichelt J., Scharte M., Schoen O., Barges M., Terol J., Climent J.,
 Navarro P., Collado C., Perez-Perez A., Ottenwaelder B., Duchemin D.,
 Cooke R., Laudie M., Berger-Liauro C., Purnelle B., Masuy D.,
 de Haan M., Maarse A.C., Alcaraz J.-P., Cottet A., Casacuberta E.,
 Monfort A., Argilou A., Flores M., Liguori R., Vitale D.,
 Mannhaupt G., Haase D., Schoof H., Rudd S., Zaccaria P., Mewes H.-W.,
 Mayer K.F.X., Kaul S., Town C.D., Koo H.L., Tallon L.J., Jenkins J.,
 Reaney T., Rizzo M., Waits A., Utterback T., Fujii C.Y., Shea T.P.,
 Creasy T.H., Haas B., Maiti R., Wu D., Peterson J., Van Aken S.,
 Pai G., Millscher J., Sellers P., Gill J.E., Feldblyum T.V.,
 Preuss D., Lin X., Nierman W.C., Salzberg S.L., White O., Venter J.C.,
 Fraser C.M., Kaneko T., Nakamura Y., Sato S., Kato T., Asamizu E.,
 Sasamoto S., Kimura T., Idesawa K., Kawashima K., Kishida Y.,
 Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
 Nakayama S., Nakazaki N., Shinozaki M., Takeuchi C., Wada T.,
 Watanabe A., Yamada M., Yasuda M., Tabata S.;
 RT "Sequence and analysis of chromosome 3 of the plant Arabidopsis
 thaliana.";
 RL Nature 408:820-822(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RA Brover V., Troukhan M., Alexandrov N., Lu Y.-P., Flavell R.,
 Feldmann K.;
 RT "Full-length cDNA from Arabidopsis thaliana.";
 RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RA Shinzaki K., Davis R.W., Ecker J.R., Theologis A.;
 RT "RIKEN Arabidopsis full length cDNA clones (RAFLs) sequenced by the
 SSP consortium (Salk/stanford/PGECL).";
 RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
 RN [7]
 RP SEQUENCE OF 36-128 FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=99311297; PubMed=10382288;
 RA Brunel D., Froger N., Pelletier G.;
 RT "Development of amplified consensus genetic markers (ACGM) in Brassica
 napus from Arabidopsis thaliana sequences of known biological
 function.";
 RL Genome 42:387-402(1999).
 RN [8]
 RP FUNCTION.
 RX PubMed=8565821;
 RA "The Arabidopsis homeotic genes APETALA3 and PISTILLATA are sufficient
 to provide the B class organ identity function.";
 RT Development 122:11-22(1996).
 RN [9]
 RP CHARACTERIZATION.
 RX PubMed=8643482;
 RA Riechmann J.L., Krizek B.A., Meyerowitz E.M.;
 RT "Dimerization specificity of Arabidopsis MADS domain homeotic proteins
 APETALA1, APETALA3, PISTILLATA, and AGAMOUS.";
 RL Proc. Natl. Acad. Sci. U.S.A. 93:4793-4798(1996).
 RN [10]
 RP GENETIC REGULATION.
 RX PubMed=11283333;
 RA Ng M., Yanofsky M.F.;
 RT "Activation of the Arabidopsis B class homeotic genes by APETALA1.";
 RL Plant Cell 13:739-753(2001).
 RN [11]
 RP CHARACTERIZATION.
 RX PubMed=11206550;
 RA Honma T., Goto K.;
 RT "Complexes of MADS-box proteins are sufficient to convert leaves into
 floral organs.";
 RL Nature 409:525-529(2001).
 CC -!- FUNCTION: Probable transcription factor involved in the genetic
 control of flower development. Is required for normal development
 of petals and stamens in the wild-type flower. Forms an
 heterodimer with PISTILLATA that is required for autoregulation of
 both AP3 and PI genes. AP3/PI heterodimer interacts with APETALA1
 or SEPALLATA3 to form a ternary complex that could be responsible
 for the regulation of the genes involved in the flower
 development.
 CC -!- SUBUNIT: Forms an heterodimer with PISTILLATA, capable of binding
 to CARG-box sequences. AP3/PI heterodimer binds AP1 or SEP3 to
 form complexes.
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- TISSUE SPECIFICITY: Expressed in petals and stamens.
 CC -!- INDUCTION: Positively regulated by the meristem identity proteins
 APETALA1 and LEAFY with the cooperation of UFO.
 CC -!- MISCELLANEOUS: Mutations in AP3 cause transformation of petals
 into sepals and stamens into carpels.
 CC -!- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSCRIPTION
 FACTORS.
 CC -!- SIMILARITY: Contains 1 K-box dimerization domain.
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 or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M86357; AAA32740.1; -;
 DR EMBL; D21125; BAA0465.1; -;
 DR EMBL; AF115798; AAD51887.1; -;
 DR EMBL; AF115799; AAD51888.1; -;

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DR EMBL; AF115800; AAD51889.1; -
DR EMBL; AF115801; AAD51890.1; -
DR EMBL; AF115802; AAD51891.1; -
DR EMBL; AF115803; AAD51892.1; -
DR EMBL; AF115804; AAD51893.1; -
DR EMBL; AF115805; AAD51894.1; -
DR EMBL; AF115806; AAD51895.1; -
DR EMBL; AF115807; AAD51896.1; -
DR EMBL; AF115808; AAD51897.1; -
DR EMBL; AF115809; AAD51898.1; -
DR EMBL; AF115810; AAD51899.1; -
DR EMBL; AF115811; AAD51900.1; -
DR EMBL; AF115812; AAD51901.1; -
DR EMBL; AF115813; AAD51902.1; -
DR EMBL; AF115814; AAD51903.1; -
DR EMBL; AL132971; CAB81799.1; -
DR EMBL; AY087369; AAM64919.1; -
DR EMBL; AY070397; AAL49893.1; -
DR EMBL; AY142590; AAN13159.1; -
DR EMBL; AF056341; AAD41557.1; -
DR FIC; A42095; A42095.
DR HSP; P11746; INNM.
DR TRANSFAC; T01776; -.
DR InterPro; IPR002487; TF_Kbox.
DR InterPro; IPR002100; TF_MADSbox.
DR Pfam; PF01486; K-box; 1.
DR Pfam; PF00319; SRP-TF; 1.
DR PRINTS; PR00404; MADSDOMAIN.
DR SMART; SM00432; MADS; 1.
DR PROSITE; PS00350; MADS_BOX_1; 1.
DR PROSITE; PS00666; MADS_BOX_2; 1.
KW Flowering; Transcription regulation; Activator; Developmental protein;
KW Nuclear protein; DNA-binding; Coiled coil; Polymorphism.
FT DOMAIN 3 37 MADS.
FT DOMAIN 93 165 K-BOX.
FT DOMAIN 75 164 COILED COIL (POTENTIAL).
FT VARIANT 31 31 K -> R (in strain cv. Lisse).
FT VARIANT 47 47 M -> T (in strain cv. Bretagne).
FT VARIANT 61 61 N -> D (in strain cv. Corsacalla-1).
FT VARIANT 73 73 T -> S (in strain cv. LI-8).
FT VARIANT 109 109 L -> V (in strain cv. Kas-1).
FT VARIANT 115 115 E -> K (in strains cv. Chi-1 and cv. Gr-3).

Query Match 35.7%; Score 51; DB 1; Length 232;
Best Local Similarity 44.4%; Pred. No. 2.3;
Matches 12; Conservative 3; Mismatches 4; Indels 8; Gaps 1;

QY 7 QRYG-----RELRRMSDEFGSK 25
Db 107 QRLGCELDLDIQLRRLEDEMENTFK 133
[1]

RESULT 11
CE05_MOUSE STANDARD; PRT; 851 AA.
AC O8K2H3;
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Protein C5orf5 homolog.
GN C5ORF5.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=FVB/N;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.N., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,

RA EMBL; Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raba S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schermer A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- SIMILARITY: Belongs to the PAM13 family.
CC -!- SIMILARITY: Contains 1 Rho-GAP domain.
CC -----
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC or send an email to license@sib-sib.ch).
CC -----
DR EMBL; BC031465; AAH31465.1; -
DR InterPro; IPR000198; RhoGAP.
DR Pfam; PF00620; RhoGAP; 1.
DR SMART; SM00324; RhoGAP; 1.
DR PROSITE; PS02338; RHO-GAP; 1.
KW GTPase activation.
FT DOMAIN 23 212 RHO-GAP.
FT DOMAIN 189 256 GLU-RICH.
SQ SEQUENCE 851 AA; 97054 MW; C2B26669FB6DB3CE CRC64;

Query Match 35.7%; Score 51; DB 1; Length 851;
Best Local Similarity 41.7%; Pred. No. 9.3;
Matches 10; Conservative 6; Mismatches 8; Indels 0; Gaps 0;

QY 1 KNLWAAQRYGRELRRMSDEFGSF 24
Db 782 EQLWKARAEKKLRKMLREFEAF 805
[1]

RESULT 12
RMUC_PSEAE STANDARD; PRT; 453 AA.
AC Q91AU3;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE DNA recombination protein rmuc homolog.
GN RMUC OR PA1031.
OS Pseudomonas aeruginosa.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=287;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 15692 / PA01;
RX MEDLINE=20437337; PubMed=10984043;
RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
RA Hickey M.J., Brinkman F.S.L., Huynh W.O., Kowalik D.J., Lagrou M.,
RA Garber R.L., Goltry L., Tolentino E., Westbrook-Wadman S., Yuan Y.,
RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
RA Smith X.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.F.,
RA Reizer J., Saler M.H., Hancock R.E.W., Lory S., Olson M.V.;
RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an
RT opportunistic pathogen.";
RN Nature 406:959-964(2000).

```

CC -!- FUNCTION: Involved in DNA recombination (By similarity).
 CC -!- SIMILARITY: BELONGS TO THE RMC FAMILY.
 CC -----
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 CC -----
 DR EMBL; AE004535; AAG04420.1; -;
 DR PIR; E83517; E83517.
 DR InterPro; IPR003798; DUF195.
 DR Pfam; PF02646; RmcC; 1.
 KW DNA recombination; Coiled coil; Complete proteome.
 FT DOMAIN 16 201 COILED COIL (POTENTIAL).
 SQ SEQUENCE 453 AA; 51539 MW; 1E7EA97E82EC5E4B CRC64;

 Query Match 34.3%; Score 49; DB 1; Length 453;
 Best Local Similarity 55.8%; Pred. No. 9.3;
 Matches 10; Conservative 4; Mismatches 2; Indels 2; Gaps 1;

 QY 4 WAAQRYGR--ELRRMSDE 19
 DB 65 WASERQGEELRLASE 82

 RESULT 13
 RAS3 RHIRA
 ID RAS3_RHIRA STANDARD; PRT; 205 AA.
 AC P22280;
 DT 01-AUG-1991 (Rel. 19, Created)
 DT 01-AUG-1991 (Rel. 19, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Ras-like protein 3.
 GN RAS3.
 OS Rhizomucor racemosus (Mucor circinelloides f. lusitanicus).
 OC Eukaryota; Fungi; Zygomycota; Zygomycetes; Mucorales; Mucoraceae;
 CC Mucor.
 CC NCBI_TaxID=4841;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 1216B;
 RX MEDLINE=91061774; PubMed=1701021;
 RA Casale W.L., McConpell D.G., Wang S.-Y., Lee Y.-J., Linz J.E.;
 RT "Expression of a gene family in the dimorphic fungus Mucor racemosus
 RL Mol. Cell. Biol. 10:6654-6663(1990).
 CC -!- ENZYME REGULATION: ALTERNATE BETWEEN AN INACTIVE FORM BOUND TO GDP
 CC AND AN ACTIVE FORM BOUND TO GTP. ACTIVATED BY A GUANINE
 CC NUCLEOTIDE-EXCHANGE FACTOR (GEF) AND INACTIVATED BY A GTPASE-
 CC ACTIVATING PROTEIN (GAP).
 CC -!- SUBCELLULAR LOCATION: Plasma membrane.
 CC -!- DEVELOPMENTAL STAGE: IN SPORULATING MYCELIUM AND MUCH LESS IN
 CC GERMLING AND YEAST.
 CC -!- SIMILARITY: BELONGS TO THE SMALL GTPASE SUPERFAMILY. RAS FAMILY.
 CC -----
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 CC -----
 DR EMBL; M55177; AAA83379.1; -;
 DR PIR; C36365; C36365.
 DR HSP; P01112; 1PL1.
 DR InterPro; IPR003577; GTPase_Ras.
 DR InterPro; IPR001806; Ras_trnsfrmg.
 DR Pfam; PF00071; ras; 1.

DR PRINTS; PR00449; RASTRNSPRMG.
 DR SMART; SM00173; RAS; 1.
 DR TIGRFAMs; TIGR00231; small_gtp; 1.
 KW GTP-Binding; Prenylation; Lipoprotein.
 FT NP_BIND 16 23 GTP (BY SIMILARITY).
 FT NP_BIND 63 67 GTP (BY SIMILARITY).
 FT NP_BIND 122 125 GTP (BY SIMILARITY).
 FT DOMAIN 38 46 EFFECTOR REGION (PROBABLE).
 FT LIPID 202 202 FARNESYL (BY SIMILARITY).
 SQ SEQUENCE 205 AA; 23408 MW; DBF086466F090F30 CRC64;

 Query Match 33.6%; Score 48; DB 1; Length 205;
 Best Local Similarity 62.5%; Pred. No. 5.6;
 Matches 10; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

 QY 11 REIRMSDEFEGRSKG 26
 DB 168 REIRRMKEQEGRSKG 183

 RESULT 14
 6PGL THEME
 ID 6PGL_THEME STANDARD; PRT; 220 AA.
 AC Q9X0N8;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 6-phosphogluconolactonase (EC 3.1.1.31) (6PGL).
 GN PGL OR DEVB OR TM1154.
 OS Thermotoga maritima.
 CC Bacteria; Thermotogae; Thermotogaceae; Thermotogales; Thermotoga.
 CC NCBI_TaxID=2336;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MSB8 / DSM 3109;
 RX MEDLINE=99287316; PubMed=10360571;
 RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
 RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
 RA McDonald L., Utterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
 RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
 RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
 RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
 RT "Evidence for lateral gene transfer between Archaea and Bacteria from
 RL genome sequence of Thermotoga maritima."
 RL Nature 399:323-329(1999).
 CC -!- FUNCTION: HYDROLYSIS OF 6-PHOSPHOGLUCONOLACTONE TO 6-
 CC PHOSPHOGLUCONATE.
 CC -!- CATALYTIC ACTIVITY: 6-phospho-D-glucono-1,5-lactone + H(2)O = 6-
 CC phospho-D-gluconate.
 CC -!- PATHWAY: pentose phosphate pathway; second step.
 CC -!- SIMILARITY: BELONGS TO THE GLUCOSAMINE/GALACTOSAMINE-6-PHOSPHATE
 CC ISOMERASE FAMILY. 6-PHOSPHOGLUCONOLACTONASE SUBFAMILY.
 CC -----
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 CC -----
 DR EMBL; AE001772; AAD36230.1; -;
 DR PIR; F72289; F72289.
 DR TIGR; TM1154; -;
 DR InterPro; IPR006148; Gluc_gal_isom.
 DR InterPro; IPR005900; Phosphogluconlac.
 DR Pfam; PF01182; Glucosamine_iso; 1.
 DR TIGRFAMs; TIGR01198; pgl; 1.
 KW Hydrolase; Complete proteome.
 SQ SEQUENCE 220 AA; 25325 MW; 950FD07E01E60C3 CRC64;

 Query Match 33.6%; Score 48; DB 1; Length 220;
 Best Local Similarity 34.8%; Pred. No. 6;

search completed: September 15, 2003, 17:00
Job time : 6 36420 sec

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 : Search time 29.3143 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-56
Perfect score: 143
Sequence: 1 KNLWAAQRYGRELRRMSDFEGSPKGL 27

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues
Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL_23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_nhc:*
- 8: sp_organelle:*
- 9: sp_phase:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	87	60.8	146	13	Q919N2
2	53	37.1	196	16	Q8VJS3
3	53	37.1	223	16	Q10843
4	53	37.1	946	11	Q8K016
5	52.5	36.7	505	8	Q47148
6	52.5	36.7	506	8	Q47149
7	52.5	36.7	506	8	Q47171
8	52.5	36.7	506	8	Q63960
9	52.5	36.7	506	8	Q62982
10	52.5	36.7	506	8	Q62975
11	52.5	36.7	506	8	Q62972
12	52.5	36.7	506	8	Q62989
13	52.5	36.7	506	8	Q62978
14	52.5	36.7	506	8	Q47155
15	52.5	36.7	506	8	Q47152
16	52.5	36.7	506	8	Q47173

17	52.5	36.7	506	8	Q62990
18	52.5	36.7	506	8	Q62974
19	52.5	36.7	506	8	Q62993
20	52.5	36.7	506	8	Q47170
21	52.5	36.7	506	8	Q47174
22	52.5	36.7	506	8	Q62983
23	52.5	36.7	506	8	Q62980
24	52.5	36.7	506	8	Q62981
25	52.5	36.7	506	8	Q62977
26	52.5	36.7	506	8	Q47168
27	52.5	36.7	506	8	Q62988
28	52.5	36.7	506	8	Q62973
29	52.5	36.7	506	8	Q62992
30	52.5	36.7	506	8	Q47175
31	52.5	36.7	506	8	Q8HSP1
32	52.5	36.7	506	8	Q8HSP0
33	52.5	36.7	506	8	Q8HSN9
34	52.5	36.7	506	8	Q8HSN8
35	52.5	36.7	506	8	Q8HSN7
36	52.5	36.7	506	8	Q8HSN6
37	52.5	36.7	506	8	Q8HSN5
38	52.5	36.7	506	8	Q8HSN4
39	52.5	36.7	507	8	Q62985
40	52.5	36.7	507	8	Q62986
41	52.5	36.7	508	8	Q62979
42	52	36.4	471	17	Q8ZY71
43	51.5	36.0	506	8	Q47153
44	51.5	36.0	506	8	Q47160
45	51.5	36.0	506	8	Q8HSP4

ALIGNMENTS

RESULT 1

Q919N2 PRELIMINARY; PRT; 146 AA.

AC Q919N2;

DT 01-OCT-2000 (TRENBLrel. 15, Created)

DT 01-DEC-2001 (TRENBLrel. 19, Last sequence update)

DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)

DE Bad.

OS Brachydanio rerio (Zebrafish) (Danio rerio).

CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

CC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;

CC Cyprinidae; Danio.

OX NCBI_TaxID=7955;

RN [1]

RP SEQUENCE FROM N.A.

RA MEDLINE=20373792; PubMed=10917738;

RA Inohara N., Nunez G.;

RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish."

RL Cell Death Differ. 7:509-510(2000).

DR EMBL; AF231017; AAF66962.2; -

DR HSP; Q92934; 1G5J.

DR ZFIN; ZDB-GENE-000616-1; bad.

SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;

Query Match 60.8%; Score 87; DB 13; Length 146;

Best Local Similarity 65.2%; Pred. No. 5.9e-05;

Matches 15; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 3 LWAQRYGRELRRMSDFEGSPK 25
|||||:|||||:|

Db 89 LWAQRYGRELRRMSDFEGSPK 111

RESULT 2

Q8VJS3 PRELIMINARY; PRT; 196 AA.

ID Q8VJS3

AC Q8VJS3;

```

Matches 10; Conservative 1; Mismatches 6; Indels 0; Gaps 0
QY      2 NLWAAQRYGRELRRMSD 18
       IIII | | | : |
DB      165 NLWAADRYNRAITARGED 181

RESULT 4
ID Q8K016 PRELIMINARY; PRT; 946 AA.
AC Q8K016;
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Inter-alpha trypsin inhibitor, heavy chain 2.
GN ITIH2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
CN NCBI_TaxId=10090;
RX [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
CA Strausberg R.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBDJ databases.
DR EMBL; BC034341; AAC34341.1; -.
DR MGD; MG1:96619; Itih2
DR InterPro; IPR006587; VIT
DR InterPro; IPR002035; VWF_A.
DR SMART; SMO0609; VIT; 1.
DR SMART; SMO0327; WVA; 1.
DR PROSITE; PS0234; VWFA; 1.
SQ SEQUENCE 946 AA; 105945 MW; 8B17DBA71B85BC5C CRC64;

Query Match 37.1%; Score 53; DB 11; Length 946;
Best Local Similarity 34.6%; Pred. No. 48;
Matches 9; Conservative 5; Mismatches 12; Indels 0; Gaps 0
QY      2 NLWAAQRYGELRRMSDEFEGSKGL 27
       I::: | | | : ||| ::||:
DB     212 NWVIMEPQGMRFLHVPDTFGHFQGV 237

RESULT 5
O47148 PRELIMINARY; PRT; 505 AA.
ID O47148
AC AC O47148;
DT 01-JUN-1998 (TREMBLrel. 06, Created)
DT 01-WAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K) (Fragment).
DN MATK.
OS Menziesia ciliicalyx.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Menziesia.
OX NCBI_TaxId=49154;
RN [1]
RP SEQUENCE FROM N.A.
RC Kron K.A.;
RL "Phylogenetics of Rhododendroideae (Ericaceae).";
RT Submitted (JUN-1996) to the EMBL/GenBank/DBDJ databases.
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
    INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
    AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
    MITOCHONDRIAL INTRONS.
DR ENBL; U61331; AAC15245.2; -.
DR InterPro; IPR000442; Intron_maturse2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.

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KW mRNA processing; Chloroplast.
PT NON_TER 1
SQ SEQUENCE 505 AA; 60233 MW; E5F927AD2B57DE5 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 505;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRLMSDEFEFGSK 25
   | : ||| : ||| | : | : |||
Db 390 KPWWAALSDSDIIERFGRIYRLNLSHYSGSLK 421

RESULT 6
O47149
ID O47149 PRELIMINARY; PRT; 506 AA.
AC O47149;
DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron kiusianum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49167;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;
RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
    INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS.
    AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
    MITOCHONDRIAL INTRONS.
DR EMBL; U61332; AAB93753.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60502 MW; 0009EA88CD28549F CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRLMSDEFEFGSK 25
   | : ||| : ||| | : | : |||
Db 391 KPWWAALSDSDIIERFGRIYRLNLSHYSGSLK 422

RESULT 7
O47171
ID O47171 PRELIMINARY; PRT; 506 AA.
AC O47171;
DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron edgeworthii.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49162;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;

```

```

RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
    INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS.
    AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
    MITOCHONDRIAL INTRONS.
DR EMBL; U61354; AAB93748.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60485 MW; 8A6353BFC5F4DC85 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRLMSDEFEFGSK 25
   | : ||| : ||| | : | : |||
Db 391 KPWWAALSDSDIIERFGRIYRLNLSHYSGSLK 422

RESULT 8
O63960
ID O63960 PRELIMINARY; PRT; 506 AA.
AC O63960;
DT 01-AUG-1998 (TREMBlrel. 07, Created)
DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MARK OR YCFL4.
OS Rhododendron tashiroi, and
    Rhododendron farrierae.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoidae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75582; /75580;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
    Yukawa I.;
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
    INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS.
    AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
    MITOCHONDRIAL INTRONS.
DR EMBL; AB012749; BAA25870.1; -.
DR EMBL; AB012745; BAA25866.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturase2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60389 MW; DE0C07AE6608E787 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

QY 1 KNLWAA-----QRYGRELRLMSDEFEFGSK 25
   | : ||| : ||| | : | : |||
Db 391 KPWWAALSDSDIIERFGRIYRLNLSHYSGSLK 422

RESULT 9
O62982
ID O62982 PRELIMINARY; PRT; 506 AA.
AC O62982;
DT 01-AUG-1998 (TREMBlrel. 07, Created)

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DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron nipponicum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75577;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matK Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
CC EMBL; AB012739; BAA25860.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60419 MW; 1F95132CCF4F6B40 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 48;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----QRYGRELRLMSDEFEGSPK 25
Db 391 KPVWAAALSDSIIEFGRIYRNLSHYSGSLK 422

RESULT 10
O62975 PRELIMINARY; PRT; 506 AA.
AC O62975;
DT 01-AUG-1998 (TREMBlrel. 07, Created)
DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron ponticum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49628;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matK Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012732; BAA25853.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60449 MW; 21DFF700B071B588 CRC64;

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Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----QRYGRELRLMSDEFEGSPK 25
Db 391 KPVWAAALSDSIIEFGRIYRNLSHYSGSLK 422

RESULT 11
O62972 PRELIMINARY; PRT; 506 AA.
AC O62972;
DT 01-AUG-1998 (TREMBlrel. 07, Created)
DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron ovatum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49169;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matK Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -!- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -!- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL; AB012729; BAA25850.1; -.
DR InterPro; IPR000442; Intron_maturase2.
DR InterPro; IPR002866; MatK_N.
DR Pfam; PF01348; Intron_maturas2; 1.
DR Pfam; PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60493 MW; D230E54B8C20FEF0 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----QRYGRELRLMSDEFEGSPK 25
Db 391 KPVWAAALSDSIIEFGRIYRNLSHYSGSLK 422

RESULT 12
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AC O62989;
DT 01-AUG-1998 (TREMBlrel. 07, Created)
DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN MATK.
OS Rhododendron indicum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=75581;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurashige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.;

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RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matk Sequences.";
RL J. Jpn. Bot. 0:0-0(1998)
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL: AB012747; BAA25868.1; -.
DR InterPro: IPR000442; Intron_maturase2.
DR InterPro: IPR002866; MatK_N.
DR Pfam: PF01348; Intron_maturas2; 1.
DR Pfam: PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60489 MW; 6D38A1D4D6FEC9BF CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----ORYGRELRMSDEFECSFK 25
Db 391 KPWAALSDDSIIEFGRIYRNLSHYSGSLK 422

RESULT 13
O62978 PRELIMINARY; PRT; 506 AA.
ID O62978 PRELIMINARY; PRT; 506 AA.
AC O62978;
DT 01-AUG-1998 (TREMBlrel. 07, Created)
DT 01-AUG-1998 (TREMBlrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN Mark.
OS Rhododendron canadense.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49465;
RN [1]
RP SEQUENCE FROM N.A.
RA Kurahige Y., Mine M., Kobayashi N., Handa T., Takayanagi K.,
RA Yukawa T.
RT "Investigation of Sectional Relationships in the Genus
RT Rhododendron(Ericaceae) based on matk Sequences.";
RL J. Jpn. Bot. 0:0-0(1998).
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL: AB012735; BAA25856.1; -.
DR InterPro: IPR000442; Intron_maturase2.
DR InterPro: IPR002866; MatK_N.
DR Pfam: PF01348; Intron_maturas2; 1.
DR Pfam: PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60350 MW; 5E832589ED64EA25 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----ORYGRELRMSDEFECSFK 25
Db 391 KPWAALSDDSIIEFGRIYRNLSHYSGSLK 422

RESULT 14
O47155 PRELIMINARY; PRT; 506 AA.
ID O47155 PRELIMINARY; PRT; 506 AA.
AC O47155;
DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN Mark.
OS Rhododendron tomentosum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49170;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;
RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL: U61335; AAB93757.1; -.
DR InterPro: IPR000442; Intron_maturase2.
DR InterPro: IPR002866; MatK_N.
DR Pfam: PF01348; Intron_maturas2; 1.
DR Pfam: PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60439 MW; 855FDBDB8A5F800D CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;

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DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN Mark.
OS Rhododendron hongkongense.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49165;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;
RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL: U61338; AAB93751.1; -.
DR InterPro: IPR000442; Intron_maturase2.
DR InterPro: IPR002866; MatK_N.
DR Pfam: PF01348; Intron_maturas2; 1.
DR Pfam: PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60406 MW; 4B5C675CE32218D8 CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;
Matches 12; Conservative 5; Mismatches 8; Indels 7; Gaps 1;

Qy 1 KNLWAA-----ORYGRELRMSDEFECSFK 25
Db 391 KPWAALSDDSIIEFGRIYRNLSHYSGSLK 422

RESULT 15
O47152 PRELIMINARY; PRT; 506 AA.
ID O47152 PRELIMINARY; PRT; 506 AA.
AC O47152;
DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Ribosomal maturase (Intron maturase) (Maturase K).
GN Mark.
OS Rhododendron tomentosum.
OG Chloroplast.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; Ericales; Ericaceae; Ericoideae; Rhodoreae; Rhododendron.
OX NCBI_TaxID=49170;
RN [1]
RP SEQUENCE FROM N.A.
RA Kron K.A.;
RT "Phylogenetics of Rhododendroideae (Ericaceae).";
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROBABLY ASSISTS IN SPLICING CHLOROPLAST GROUP II
CC INTRONS (BY SIMILARITY).
CC -1- SIMILARITY: WITH CORRESPONDING ORF IN OTHER PLANT CHLOROPLASTS,
CC AND REGIONS OF SIMILARITY TO MATURASE-LIKE POLYPEPTIDES ENCODED BY
CC MITOCHONDRIAL INTRONS.
DR EMBL: U61335; AAB93757.1; -.
DR InterPro: IPR000442; Intron_maturase2.
DR InterPro: IPR002866; MatK_N.
DR Pfam: PF01348; Intron_maturas2; 1.
DR Pfam: PF01824; MatK_N; 1.
KW mRNA processing; Chloroplast.
SQ SEQUENCE 506 AA; 60439 MW; 855FDBDB8A5F800D CRC64;

Query Match 36.7%; Score 52.5; DB 8; Length 506;
Best Local Similarity 37.5%; Pred. No. 28;

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D6	

Search completed: September 15, 2003, 17:25:50
Job time : 29.3143 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:01 : Search time 22.6286 Seconds
(without alignments)
112.231 Million cell updates/sec

Title: US-09-544-664-28

Perfect score: 84

Sequence: 1 QRYGRLRMSDEPEG (16)

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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A_Geneseq_13Jun03.*

- 1: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1980.DAT.*
- 2: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1981.DAT.*
- 3: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1982.DAT.*
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- 5: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1984.DAT.*
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- 24: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	84	100.0	16	20 AAY05422	Mouse BAD BH3 doma
2	84	100.0	16	21 AAB37028	Bcl2 polypeptide B
3	84	100.0	23	17 AAR95156	bcl-x(L)/bcl-2 ass
4	84	100.0	26	21 AAB37001	Bcl2 polypeptide B
5	84	100.0	26	21 AAB37002	Bcl2 polypeptide B
6	84	100.0	27	21 AAB37003	Bcl2 polypeptide B
7	84	100.0	27	21 AAB37056	Bcl2 polypeptide B
8	84	100.0	28	21 AAB37055	Bcl2 polypeptide B
9	84	100.0	162	22 AAB70370	Shorter murine BAD

10	84	100.0	204	17 AAR95168	bcl-x(L)/bcl-2 ass
11	84	100.0	204	19 AAW61315	Murine BCL-XL/BCL-
12	84	100.0	204	19 AAW61316	Mutant BCL-XL/BCL-
13	84	100.0	204	19 AAW61317	Mutant BCL-XL/BCL-
14	84	100.0	204	19 AAW61318	Mutant BCL-XL/BCL-
15	84	100.0	204	19 AAW58832	Murine BAD protein
16	84	100.0	204	22 AAB70369	Longer murine BAD
17	84	100.0	204	24 ABR39082	Murine BAD protein
18	84	100.0	567	22 AAU00220	Bad-DRTR apoptosis
19	73	86.9	16	20 AAY05421	Human BAD BH3 doma
20	73	86.9	16	21 AAB37029	Bcl2 polypeptide B
21	73	86.9	18	23 AAB78483	Human Bcl2 fluores
22	73	86.9	18	23 AAB78609	Human Bcl2 peptide
23	73	86.9	20	23 AAB78459	Mutant Bcl2 compet
24	73	86.9	20	23 AAB78626	Human Bad peptide
25	73	86.9	21	23 AAB78500	Mutant Bcl2 compet
26	73	86.9	21	23 AAB78608	Human Bad peptide
27	73	86.9	21	23 AAB78630	Human Bad peptide
28	73	86.9	22	23 AAB78629	Human Bad peptide
29	73	86.9	22	23 AAB78628	Human Bad peptide
30	73	86.9	24	23 AAB78482	Human Bcl2 fluores
31	73	86.9	24	23 AAB78605	Human Bad peptide
32	73	86.9	24	23 AAB78627	Human Bad peptide
33	73	86.9	25	23 AAB56161	PTPC-interacting T
34	73	86.9	25	23 AAB78481	Human Bcl2 fluores
35	73	86.9	25	23 AAB78484	Mutant Bcl2 compet
36	73	86.9	25	23 AAB78485	Mutant Bcl2 compet
37	73	86.9	25	23 AAB78486	Mutant Bcl2 compet
38	73	86.9	25	23 AAB78487	Mutant Bcl2 compet
39	73	86.9	25	23 AAB78488	Mutant Bcl2 compet
40	73	86.9	25	23 AAB78489	Mutant Bcl2 compet
41	73	86.9	25	23 AAB78490	Mutant Bcl2 compet
42	73	86.9	25	23 AAB78491	Mutant Bcl2 compet
43	73	86.9	25	23 AAB78492	Mutant Bcl2 compet
44	73	86.9	25	23 AAB78493	Mutant Bcl2 compet
45	73	86.9	25	23 AAB78494	Mutant Bcl2 compet

ALIGNMENTS

RESULT 1

AAV05422

ID AAY05422 standard; peptide; 16 AA.

XX AAY05422;

02-JUL-1999 (first entry)

DE Mouse BAD BH3 domain.

KW BH3 domain; cell death agonist; bcl homology domain; BCL-2 family;
KW apoptosis promoter; cancer cell; virus infected cell; inflammation;
KW autoantibody producing cell; cancer; lymphoproliferative condition;
KW arthritis; autoimmune disease; therapy.

OS Mus sp.

PN WO9916787-A1.

08-APR-1999

22-SEP-1998; 98WO-US19765.

PR 07-OCT-1997; 97US-0946039.

PR 26-SEP-1997; 97US-0060133.

PA (UNIV) UNIV WASHINGTON.

PI Korsmeyer SJ;

XX WPI; 1999-255058/21.

10261 (1 Aug 1999)

PT Bcl homology domain 3 polypeptide
XX
PS Example 1; Fig 4; 104pp; English.

XX
CC This sequence represents the BH3 domain of mouse BAD.
CC The invention relates to a bcl homology domain 3 (BH3 domain),
CC derived from a proapoptotic member of the BCL-2 family. The
CC BH3 polypeptide can be used in a method for promoting apoptosis in a
CC target cell, especially where the cell is a cancer cell a virus infected
CC cell or an autotantibody producing cell. The BH3 polypeptide can be used
CC in therapeutic compositions for treating disease including cancer, other
CC lymphoproliferative conditions, arthritis, inflammation, and autoimmune
CC diseases, which may result from the down regulation of cell death
XX regulation.

XX Sequence 16 AA;

Query Match 100.0%; Score 84; DB 20; Length 16;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRLMSDFEG 16
Db 1 QRYGRELRLMSDFEG 16

RESULT 2
AAB37028
ID AAB37028 standard; peptide; 16 AA.
XX
AC AAB37028;
XX
DT 28-FEB-2001 (first entry)
XX
DE Bcl2 polypeptide BH3 domain peptide #28.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.

XX Homo sapiens.

XX WO2000059526-A1.

XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UJJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer -

PS Claim 18; Page 18; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH₂ or OH; or X = O or NH,
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH₂; and R = 2-18C alkyl or alkoxy/ 2-14C alkyl/enyl containing one
CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally

CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the peptide portion of the conjugate. The peptides represent analogues
CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX Sequence 16 AA;

Query Match 100.0%; Score 84; DB 21; Length 16;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRLMSDFEG 16
Db 1 QRYGRELRLMSDFEG 16

RESULT 3
AAR95166
ID AAR95166 standard; peptide; 23 AA.

XX AAR95166;

XX 03-JAN-1997 (first entry)

XX bcl-x(L)/bcl-2 associated death promoter epitope, residues 138-160.

XX Epitope: murine; bcl-x(L)/bcl-2 associated death promoter; Bad; stroke;
KW polypeptide; bcl-x; cell death; regulate; BHL; BH2; apoptotic cell death;
KW cytokine deprivation; IL-3 dependent cell line; immunodeficiency; AIDS;
KW neurodegenerative disease; senescence; ischaemia; neoplasia.

XX Mus musculus.

XX WO9613614-A1.

XX 09-MAY-1996.

XX 31-OCT-1995; 95WO-US14246.

XX 31-OCT-1994; 94US-0333565.

XX (UNIW) UNIV WASHINGTON.

XX Korsmeyer SJ;

XX WPI; 1996-251465/25.

XX Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
PT useful to treat neoplasia and apoptosis and to identify agents
PT inhibiting its binding to bcl-2 or bcl-x(L) to form heteromultimers

XX Claim 2; Page 103; 130pp; English.

XX The sequences given in AAR95155-67 represent epitopes derived from the
CC murine bcl-x(L)/bcl-2 associated death promoter (Bad) polypeptide (see
CC also AAR95168). Bad is a 22.1 kD protein which interacts with bcl-2 and
CC bcl-x proteins and regulates cell death. It has homology to the bcl-2-
CC related family clustered in the BHL and BH2 domain. Bad has been found
CC to hybridise to bcl-x(L) and bcl-2 in yeast two-hybrid assays and in
CC vivo in mammalian cells. Overexpressed Bad counters the death

CC inhibitory activity of bcl-x(L), but is much less effective at countering
 CC the death inhibitory activity of bcl-2. Bad expression can accelerate
 CC apoptotic cell death induced by cytokine deprivation in an IL-3 dependent
 CC cell line expressing bcl-x(L), and its also counters the death repressor
 CC activity of bcl-x(L). Bad competes with Bax for binding to bcl-x(L).
 CC Bad may be used to identify agents which inhibit its binding to bcl-2
 CC or bcl-x(L) to form heterodimers. Such agents may be used to treat
 CC neurodegenerative diseases, immunodeficiency diseases, e.g. AIDS,
 CC senescence or ischaemia.

XX Sequence 23 AA;

Query Match 100.0%; Score 84; DB 17; Length 23;
 Best Local Similarity 100.0%; Pred. No. 3.7e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDFEG 16
 |||||
 Db 8 QRYGRELRRMSDFEG 23

RESULT 4
 AAB37001
 ID AAB37001 standard; peptide; 26 AA.

XX AAB37001;
 XX
 XX 28-FEB-2001 (first entry)
 XX
 XX Bcl2 polypeptide BH3 domain peptide #1.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.

XX WC2000059526-AL.

XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UYJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer

XX Claim 18; Page 17; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of

CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX Sequence 26 AA;

Query Match 100.0%; Score 84; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 4.2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDFEG 16
 |||||
 Db 6 QRYGRELRRMSDFEG 21

RESULT 5
 AAB37002
 ID AAB37002 standard; peptide; 26 AA.

XX AAB37002;

XX 28-FEB-2001 (first entry)

XX Bcl2 polypeptide BH3 domain peptide #2.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.

XX Homo sapiens.

XX WC2000059526-AL.

XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UYJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer

XX Claim 18; Page 17; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
 CC (R-X)n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one
 CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain

CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
 XX
 SQ Sequence 26 AA;
 Query Match 100.0%; Score 84; DB 21; Length 26;
 Best Local Similarity 100.0%; Pred. No. 4.2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 Db 6 QRYGRELRLMSDFEG 21
 RESULT 6
 AAB37003
 ID AAB37003 standard; peptide; 27 AA.
 XX
 AC AAB37003;
 XX
 DT 28-FEB-2001 (first entry)
 XX
 DE Bcl2 polypeptide BH3 domain peptide #3.
 XX
 KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.
 XX
 OS Homo sapiens.
 XX
 PN WO200059526-A1.
 XX
 PD 12-OCT-2000.
 XX
 PF 06-APR-2000; 2000WO-US09352.
 XX
 PR 07-APR-1999; 99US-0128202.
 XX
 PA (UYJE-) UNIV JEFFERSON THOMAS.
 XX
 PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX
 DR WPI; 2000-679325/66.
 XX
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 XX
 PS Claim 18; Page 17; 74pp; English.
 XX
 CC The invention relates to a peptide conjugate having the formula:
 CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one

CC or two double bonds, cyclobutyl, cyclopentyl, cyclohexyl optionally
 CC monosubstituted with a 1-5C straight or branched chain alkyl group,
 CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
 CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
 CC of the peptide portion of the conjugate. The peptides represent analogues
 CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
 CC the BH3 domain of the cell death agonist Bad. The peptide conjugate is
 CC useful for modulating apoptosis in the cells of a subject, or for
 CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
 CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
 CC function. In particular, the peptide conjugate is useful for treating a
 CC subject afflicted with a cancer characterized by cancer cells that
 CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
 CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
 CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
 CC conjugate is also useful for treating disorders characterized by
 CC increased apoptosis, e.g. neurodegenerative disorders, acquired
 CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.
 XX
 SQ Sequence 27 AA;
 Query Match 100.0%; Score 84; DB 21; Length 27;
 Best Local Similarity 100.0%; Pred. No. 4.4e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 Db 6 QRYGRELRLMSDFEG 21
 RESULT 7
 AAB37056
 ID AAB37056 standard; peptide; 27 AA.
 XX
 AC AAB37056;
 XX
 DT 28-FEB-2001 (first entry)
 XX
 DE Bcl2 polypeptide BH3 domain peptide #56.
 XX
 KW Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
 KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; Bad;
 KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
 KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
 KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
 KW stroke; myocardial infarction.
 XX
 OS Homo sapiens.
 XX
 PN WO200059526-A1.
 XX
 PD 12-OCT-2000.
 XX
 PF 06-APR-2000; 2000WO-US09352.
 XX
 PR 07-APR-1999; 99US-0128202.
 XX
 PA (UYJE-) UNIV JEFFERSON THOMAS.
 XX
 PI Huang Z, Wang J, Zhang Z, Shan S, Lu Z;
 XX
 DR WPI; 2000-679325/66.
 XX
 XX New peptide conjugates for modulating apoptosis or for inhibiting B
 PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
 PT treating neurodegenerative disorders, stroke, or cancer -
 XX
 PS Claim 18; Page 19; 74pp; English.
 XX
 CC The invention relates to a peptide conjugate having the formula:
 CC (R-X)_n-peptide where n = 1-10; X = C=O, when the R-X group is attached
 CC to the N-terminus of the peptide, or a side chain of the peptide where
 CC the functional group of the side chain is NH2 or OH; or X = O or NH,
 CC when the R-X group is attached to the C-terminus of the peptide, or a
 CC side chain of the peptide, where the side chain functional group is COOH
 CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylenyl containing one

PT apoptosis, comprises amino acid substitutions at Ser118, Ser155 or
 XX Ser113 -
 PS Claim 7; Page 148-149; 157pp; English.
 XX
 CC The present invention describes an isolated or synthetic polypeptide
 CC (1) comprising a less than full length amino acid sequence of a mutant
 CC Bcl-XL/Bcl-2 associated cell death regulator polypeptide (BAD) or its
 CC fragment, which contains amino acid substitutions at Ser118 of a human
 CC BAD, Ser155 of a murine BAD (longer murine BAD) or Ser113 of a murine
 CC BAD (shorter murine BAD). (1) has immunostimulant, neuroprotective,
 CC antitoxic, antischismatic, vulnerary, cytoskeletal, antiviral,
 CC neurotrophic, antiischemic, antiinflammatory and immunosuppressive activities, and
 CC can be used as an apoptosis inducer or inhibitor. BAD polypeptides and
 CC polynucleotides can be used for screening candidate compounds and drugs
 CC for activity that promote cell survival or apoptosis. Other uses include
 CC inducing or inhibiting apoptosis in a cell. Candidate compounds
 CC identified and (mutant) BAD polypeptides are useful in treating
 CC immunodeficiency diseases, neurodegenerative diseases, ischaemic cell
 CC death, reperfusion cell death, wound healing, cancer, viral infections,
 CC lymphoproliferative conditions, arthritis, infertility, inflammation and
 CC autoimmune diseases. The present sequence represents a specifically
 CC claimed shorter murine BAD mutant amino acid sequence from the present
 CC invention.
 XX Sequence 162 AA;
 SQ
 Query Match 100.0%; Score 84; DB 22; Length 162;
 Best Local Similarity 100.0%; Pred. No. 2.7e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 Db 103 QRYGRELRLMSDFEG 118
 |||||
 RESULT 10
 AAR95168
 ID AAR95168 standard; Protein; 204 AA.
 XX AAR95168;
 AC
 XX 06-JAN-1997 (first entry)
 DT
 XX
 DE bcl-x(L)/bcl-2 associated death promoter protein.
 XX
 KW Epitope; murine; bcl-x(L)/bcl-2 associated death promoter; Bad; stroke;
 KW polypeptide; bcl-x; cell death; regulate; BH1; BH2; apoptotic cell death;
 KW cytokine deprivation; IL-3 dependent cell line; immunodeficiency; AIDS;
 KW neurodegenerative disease; senescence; ischaemia; neoplasia.
 XX
 OS Mus musculus.
 XX
 FH Key Location/Qualifiers
 FT Region 147..149
 FT /note= "BH1 conserved amino acids"
 FT Region 191..192
 FT /note= "BH2 conserved amino acids"
 FT Domain 38..61
 FT /note= "PEST sequence"
 FT Domain 111..130
 FT /note= "PEST sequence"
 XX
 XX W09613614-AL.
 PN
 XX 09-MAY-1996.
 PD
 XX 31-OCT-1995; 95WO-US14246.
 XX
 XX 31-OCT-1994; 94US-0333565.
 PR
 XX (UNIW) UNIV WASHINGTON.
 PA
 XX Claim 1; Fig 10; 95pp; English.

PI Korsmeyer SJ;
 XX
 DR WPI; 1996-251465/25.
 DR N-PSDB; AAT29479.
 XX
 PT Polynucleotide encoding bcl-x(L)/bcl-2 associated death promoter -
 PT useful to treat neoplasia and apoptosis and to identify agents
 PT inhibiting its binding to bcl-2 or bcl-x(L) to form heteromultimers
 XX
 PS Claim 3; Fig 1; 130pp; English.
 XX
 CC This sequence represents the murine bcl-x(L)/bcl-2 associated death
 CC promoter (Bad) gene. Bad is a 22.1 kD protein which interacts with
 CC bcl-2 and bcl-x proteins and regulates cell death. It has homology
 CC to the bcl-2-related family clustered in the BH1 and BH2 domain. Bad
 CC has been found to hybridize to bcl-x(L) and bcl-2 in yeast two-hybrid
 CC assays and in vivo in mammalian cells. Overexpressed Bad counters the
 CC death inhibitory activity of bcl-x(L), but is much less effective at
 CC countering the death inhibitory activity of bcl-2. Bad expression can
 CC accelerate apoptotic cell death induced by cytokine deprivation in an
 CC IL-3 dependent cell line expressing bcl-x(L), and its also counters the
 CC death repressor activity of bcl-x(L). Bad competes with Bax for binding
 CC to bcl-x(L). Bad may be used to identify agents which inhibit its
 CC binding to bcl-2 or bcl-x(L) to form heterodimers. Such agents may be
 CC used to treat neurodegenerative diseases, immunodeficiency diseases,
 CC e.g. AIDS, senescence or ischaemia.
 XX
 SQ Sequence 204 AA;
 Query Match 100.0%; Score 84; DB 17; Length 204;
 Best Local Similarity 100.0%; Pred. No. 3.5e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 Db 145 QRYGRELRLMSDFEG 160
 |||||
 RESULT 11
 AAW61315
 ID AAW61315 standard; Protein; 204 AA.
 XX
 AC AAW61315;
 XX
 DT 07-OCT-1998 (first entry)
 DE
 DE Murine BCL-XL/BCL-2 associated cell death regulator.
 XX
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX
 OS Mus sp.
 XX
 XX W09817682-AL.
 PN
 XX 30-APR-1998.
 PD
 XX 17-OCT-1997; 97WO-US19175.
 PF
 XX 18-OCT-1996; 96US-0733505.
 PR
 XX (UNIW) UNIV WASHINGTON.
 PA
 XX Korsmeyer SJ;
 PI
 XX WPI; 1998-261422/23.
 DR
 DR N-PSDB; AAV27833.
 XX
 PT New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 XX
 XX Claim 1; Fig 10; 95pp; English.

XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence is the murine BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX
 XX Sequence 204 AA;
 SQ

Query Match 100.0%; Score 84; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. NO. 3.5e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRLRMSDEFEG 16
 |||||||||||||||
 Db 145 QRYGRELRLRMSDEFEG 160

RESULT 12
 AAW61316
 ID AAW61316 standard; Protein; 204 AA.
 XX
 AC AAW61316;
 XX
 XX 07-OCT-1998 (first entry)
 DT
 DT
 DE Mutant BCL-XL/BCL-2 associated cell death regulator #1.
 XX
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 KW
 XX Mus sp.
 OS Synthetic.
 OS
 XX WO9817682-A1.
 PN
 XX 30-APR-1998.
 PD
 XX 17-OCT-1997; 97WO-US19175.
 PF
 XX 18-OCT-1996; 96US-0733505.
 PR
 XX (UNIW) UNIV WASHINGTON.
 PA
 XX Korsmeyer SJ;
 PI
 XX WPI; 1998-261422/23.
 DR
 DR N-PSDB; AAV27834.
 XX
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 PT
 XX Claim 7; Page 59; 95pp; English.
 PS
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell

CC The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.
 XX
 XX Sequence 204 AA;
 SQ

Query Match 100.0%; Score 84; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. NO. 3.5e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRLRMSDEFEG 16
 |||||||||||||||
 Db 145 QRYGRELRLRMSDEFEG 160

RESULT 13
 AAW61317
 ID AAW61317 standard; Protein; 204 AA.
 XX
 AC AAW61317;
 XX
 XX 07-OCT-1998 (first entry)
 DT
 DT
 DE Mutant BCL-XL/BCL-2 associated cell death regulator #2.
 XX
 XX Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 KW serine substituted mutant; apoptosis; cancer; viral infection.
 KW
 XX Mus sp.
 OS Synthetic.
 OS
 XX WO9817682-A1.
 PN
 XX 30-APR-1998.
 PD
 XX 17-OCT-1997; 97WO-US19175.
 PF
 XX 18-OCT-1996; 96US-0733505.
 PR
 XX (UNIW) UNIV WASHINGTON.
 PA
 XX Korsmeyer SJ;
 PI
 XX WPI; 1998-261422/23.
 DR
 DR N-PSDB; AAV27835.
 XX
 XX New mutant BAD polypeptide with phosphorylatable serine replaced -
 PT useful for, e.g. treating reduced apoptosis such as in cancer or
 PT viral infection
 PT
 XX Claim 7; Page 60; 95pp; English.
 PS
 XX The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell

CC death regulator) proteins, having an amino acid other than Ser at
 CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.

XX SQ Sequence 204 AA;
 Query Match 100.0%; Score 84; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 3.5e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 |||||
 Db 145 QRYGRELRLMSDFEG 160

RESULT 14
 AAW61318
 ID AAW61318 standard; Protein; 204 AA.
 XX AC AAW61318;
 XX DT 07-OCT-1998 (first entry)
 XX DE Mutant BCL-XL/BCL-2 associated cell death regulator #3.
 XX KW Murine; mouse; BCL-XL/BCL-2 associated cell death regulator; BAD protein;
 XX KW serine substituted mutant; apoptosis; cancer; viral infection.
 XX OS Mus sp.
 XX OS Synthetic.
 XX PN WO9817682-A1.
 XX PD 30-APR-1998.
 XX PF 17-OCT-1997; 97WO-US19175.
 XX PR 18-OCT-1996; 96OS-0733505.
 XX PA (UNIW) UNIV WASHINGTON.
 XX PI Korsmeyer SJ;
 XX DR WPI; 1998-261422/23.
 XX DR N-PSDB; AAV27836.
 XX PT New mutant BAD polypeptide with phosphorylatable serine replaced -
 XX PT useful for, e.g. treating reduced apoptosis such as in cancer or
 XX PT viral infection
 XX PS Claim 7; Page 60-61; 95pp; English.
 XX CC The present invention describes mutant BAD (BCL-XL/BCL-2 associated cell
 CC death regulator) proteins, having an amino acid other than Ser at

CC position 112 and/or 136, relative to the murine BAD 204 aa sequence. The
 CC present sequence represents a mutant BAD protein. Also described are: (1)
 CC fragments of mutant BAD protein able to decrease cell viability; (2)
 CC fusion proteins of mutant BAD with a heterologous polypeptide that
 CC increases intracellular delivery. Mutant BAD proteins are used to treat
 CC or prevent diseases associated with reduced apoptosis, e.g. cancer,
 CC viral infection, lymphoproliferation, arthritis, infertility,
 CC inflammation and autoimmune disease. Polynucleotide sequences encoding
 CC mutant BAD proteins can be used similarly by gene therapy or to produce
 CC transgenic animals for use as disease models or in drug screening. BAD
 CC proteins phosphorylated at specified Ser are used to screen for enhancers
 CC and inhibitors of serine-phosphatase. Inhibitors are potentially useful
 CC in treatment of excessive apoptosis such as AIDS, neurodegeneration,
 CC aging or ischaemic cell death. The apoptotic status of cells is
 CC determined by measuring relative amounts of phosphorylated and non-
 CC phosphorylated BAD, by usual immunoassays. Mutant BAD proteins have
 CC greater death-promoting activity than wild-type BAD which can become
 CC phosphorylated on the specified Ser, forming a product that does not
 CC heterodimerise with BCL-2 or BCL-XL but instead binds to 14-3-3 family
 CC proteins in the cytosol, thus promoting cell survival. The mutants with
 CC Ser substituted cannot bind 14-3-3.

XX SQ Sequence 204 AA;
 Query Match 100.0%; Score 84; DB 19; Length 204;
 Best Local Similarity 100.0%; Pred. No. 3.5e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRLMSDFEG 16
 |||||
 Db 145 QRYGRELRLMSDFEG 160

RESULT 15
 AAW58832
 ID AAW58832 standard; Protein; 204 AA.
 XX AC AAW58832;
 XX DT 23-JUL-1998 (first entry)
 XX DE Murine BAD protein.
 XX KW BAD protein; BCL-XL/BCL-2 associated cell death regulator; 14-3-3;
 XX KW serine phosphorylation; post-translational modification; apoptosis;
 XX KW signal transduction regulator; phosphoserine phosphatase; senescence;
 XX KW immunodeficiency disease; neurodegenerative disease; infertility;
 XX KW cancer; viral infection; lymphoproliferative condition; arthritis;
 XX KW inflammation; autoimmune diseases.
 XX OS Mus sp.
 XX PN WO9809643-A1.
 XX PD 12-MAR-1998.
 XX PF 09-SEP-1997; 97WO-US15871.
 XX PR 09-SEP-1996; 96US-0707868.
 XX PA (UNIW) UNIV WASHINGTON.
 XX PI Korsmeyer SJ;
 XX DR WPI; 1998-207049/18.
 XX PT Serine-phosphorylated Bcl-X-L/Bcl-2 Associated cell Death regulator
 XX PT polypeptide - useful for modulation of apoptosis associated with,
 XX PT e.g. cancer and immunodeficiency diseases
 XX PS Claim 3; Fig 8; 61pp; English.
 XX CC This sequence represents a novel serine-phosphorylated protein, BAD


```
RESULT 2
US-08-661-479-10
; Sequence 10, Application US/08661479
; Patent No. 5834209
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/561.479
; FILING DATE: 11-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 23 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-08-661-479-10

Query Match 100.0%; Score 84; DB 2; Length 23;
Best Local Similarity 100.0%; Pred. No. 5.4e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRMSDEFG 16
Db 8 QRYGRELRMSDEFG 23
|||||

RESULT 3
US-08-333-565-2
; Sequence 2, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/661.479
; FILING DATE: 11-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 23 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-08-661-479-10

Query Match 100.0%; Score 84; DB 2; Length 23;
Best Local Similarity 100.0%; Pred. No. 5.4e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRMSDEFG 16
Db 8 QRYGRELRMSDEFG 23
|||||
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```
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/333,565
FILING DATE: 31-OCT-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 15726A-000700
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 326-2400
TELEFAX: (415) 326-2422
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
FEATURE:
NAME/KEY: Protein
LOCATION: 1..204
OTHER INFORMATION: /note= "Deduced amino acid sequence
of mouse Bcl-x."
US-08-333-565-2

Query Match 100.0%; Score 84; DB 1; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRMSDEFG 16
Db 145 QRYGRELRMSDEFG 160
|||||

RESULT 4
US-08-661-479-2
; Sequence 2, Application US/08661479
; Patent No. 5834209
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; TITLE OF INVENTION: REGULATOR
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Khourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/661.479
; FILING DATE: 11-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
```

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; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..204
; OTHER INFORMATION: /note= "deduced amino acid sequence
; OTHER INFORMATION: of mouse BAD."
US-08-661-479-2

Query Match 100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRLMSDFEG 16
   |||||
Db 145 QRYGRELRLMSDFEG 160

RESULT 5
US-08-733-505A-1
; Sequence 1, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-733-505A-1

Query Match 100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRLMSDFEG 16
   |||||
Db 145 QRYGRELRLMSDFEG 160

RESULT 6
US-08-733-505A-12
; Sequence 12, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-733-505A-12

Query Match 100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRLMSDFEG 16
   |||||
Db 145 QRYGRELRLMSDFEG 160

RESULT 7
US-08-733-505A-13
; Sequence 13, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-08-733-505A-13

Query Match 100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRLMSDFEG 16
   |||||
Db 145 QRYGRELRLMSDFEG 160
```

```

; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-08-733-505A-13

Query Match      100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEFG 16
Db      145 QRYGRLRMSDEFG 160

RESULT 8
; US-08-733-505A-14
; Sequence 14, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-08-733-505A-14

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Query Match      100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 5.5e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEFG 16
Db      145 QRYGRLRMSDEFG 160

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RESULT 9
; US-08-717-123-3
; Sequence 3, Application US/08717123
; Patent No. 5965703
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, Encoding Nucleic
; TITLE OF INVENTION: ACIDS and Methods of Use
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/717,123
; FILING DATE: 20-SEP-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-ID 1929
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 204 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; US-08-717-123-3

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Query Match      96.4%; Score 81; DB 2; Length 204;
Best Local Similarity 93.8%; Pred. No. 1.8e-06;
Matches 15; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEFG 16
Db      145 QRYGRLRMSDEFG 160

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RESULT 10
; US-09-375-257-3
; Sequence 3, Application US/09375257
; Patent No. 6504022
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.42801
; CURRENT APPLICATION NUMBER: US/09/375,257
; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
; US-09-375-257-3

```

```

Query Match      96.4%; Score 81; DB 4; Length 204;
Best Local Similarity 93.8%; Pred. No. 1.8e-06;
Matches 15; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

```

```
QY      1 QRYGRELRRMSDEFEG 16
        |||||:||||
Db     145 QRYGRELRRNTDEFEG 160
```

RESULT 11

```

US-08-733-505A-55
; Sequence 55, Application US/08733505A
; Patent No. 5856445
;
; GENERAL INFORMATION:
;
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSER: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSITH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: FLOPPY disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:

```

```

/ CLASSIFICATION: 30
/
/ ATTORNEY/AGENT INFORMATION:
/
/   NAME:  HOLLAND, DONALD R.
/
/   REGISTRATION NUMBER:  35,197
/
/   REFERENCE/DOCKET NUMBER:  965458
/
/ TELECOMMUNICATION INFORMATION:
/
/   TELEPHONE:  (314) 427-5188
/
/   TELEFAX:  (314) 727-6092
/
/   INFORMATION FOR SEQ ID NO:  55:
/
/     SEQUENCE CHARACTERISTICS:
/
/       LENGTH:  59 amino acids
/
/       TYPE:  amino acid
/
/       STRANDEDNESS:
/
/         TOPOLOGY:  linear
/
/         MOLECULE TYPE:  peptide
/
/   PS-08-733-505A-55

```

```
Query Match      86.9%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 QRYGRELRRMSDEF 14
        |||||
Db      46 QRYGRELRRMSDEF 59
```

RESULT 12

US-08-733-505A-56
; Sequence 56, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:

```

1 MEDIUM TYPE: Floppy disk
2
3 COMPUTER: IBM PC compatible
4
5 OPERATING SYSTEM: PC-DOS/MS-DOS
6
7 SOFTWARE: PatentIn Release #1.0, Version #1.30
8
9 CURRENT APPLICATION DATA:
10
11 APPLICATION NUMBER: US/08/733,505A
12

```

```

CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 5
SEQUENCE CHARACTERISTICS:
LENGTH: 59 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-56

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Query Match 86.9%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels

QY 1 QRYGRELRRMSDEF 14
db 46 QRYGRELRRMSDEF 59

RESULT 13

US-06-733-505A-57
; Sequence 57, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105

```

:
:
: COMPUTER READABLE FORM:
:
: MEDIUM TYPE: Floppy disk
:
: COMPUTER: IBM PC compatible
:
: OPERATING SYSTEM: PC-DOS/MS-DOS
:
: SOFTWARE: Patentin Release #1.0, Version #1.30
:
: CURRENT APPLICATION DATA:
:
: APPLICATION NUMBER: US/08/733.505A
:
:
:

```

```

/ CLASSIFICATION: 530
/
/ ATTORNEY/AGENT INFORMATION:
/ NAME: HOLLAND, DONALD R.
/
/ REGISTRATION NUMBER: 35,197
/
/ REFERENCE/DOCKET NUMBER: 965458
/
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (314) 727-5188
/ TELEFAX: (314) 727-6092
/
/ INFORMATION FOR SEQ ID NO: 57:
/
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 59 amino acids
/ TYPE: amino acid
/ STRANDEDNESS:
/
/ TOPOLOGY: linear
/
/ MOLECULE TYPE: peptide
/
/ US-08-733-515A-57

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Query Match 86.9%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRMSDEF 14
Db 46 QRYGRELRMSDEF 59

RESULT 14

US-08-733-505A-58
; Sequence 58, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 58:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-58

Query Match 86.9%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRMSDEF 14
Db 46 QRYGRELRMSDEF 59

RESULT 15

US-08-665-617-2
; Sequence 2, Application US/08665617
; Patent No. 566316
; GENERAL INFORMATION:
; APPLICANT: Xudong, Yin
; TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Saliwanchik & Saliwanchik
; STREET: 2421 N.W. 41st Street, Suite A-1
; CITY: Gainesville
; STATE: Florida

COUNTRY: USA
; ZIP: 32606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/665,617
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Saliwanchik, David R.
; REGISTRATION NUMBER: 31,794
; REFERENCE/DOCKET NUMBER: CL-8
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (352) 375-8100
; TELEFAX: (352) 372-5800
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 166 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-665-617-2

Query Match 86.9%; Score 73; DB 1; Length 166;
Best Local Similarity 100.0%; Pred. No. 3.4e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRMSDEF 14
Db 106 QRYGRELRMSDEF 119

Search completed: September 15, 2003, 17:45:06
Job time : 9.34286 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:25:56 ; Search time 12.6857 Seconds
(without alignments)
184.034 Million cell updates/sec

Title: US-09-544-664-28

Perfect score: 84

Sequence: 1 QRYGRLRRMSDEFEQ 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 541936 seqs, 145912426 residues

Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Published_Applications_AA.*

1: /cgn2_6/ptodata/1/pubaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubaa/PCTUS_PUBCOMB.pep.*
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9: /cgn2_6/ptodata/1/pubaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	81	96.4	204	9 US-09-922-378-3	Sequence 3, Appli
2	81	96.4	204	14 US-10-066-179-3	Sequence 3, Appli
3	73	86.9	25	15 US-10-059-261-258	Sequence 258, App
4	73	86.9	168	9 US-09-922-378-2	Sequence 2, Appli
5	73	86.9	168	9 US-09-894-657-1	Sequence 1, Appli
6	73	86.9	168	9 US-09-894-657-7	Sequence 7, Appli
7	73	86.9	168	14 US-10-066-179-2	Sequence 2, Appli
8	67	79.8	15	15 US-10-174-105A-147	Sequence 147, App
9	50	59.5	215	15 US-10-156-761-9145	Sequence 9145, Ap
10	43	51.2	213	9 US-09-843-846-18	Sequence 18, Appl
11	41	48.8	582	10 US-09-331-631A-22	Sequence 22, Appl
12	40	47.6	380	15 US-10-156-761-13594	Sequence 13594, A
13	40	47.6	2871	15 US-10-146-473-41	Sequence 41, Appl
14	38.5	45.8	764	15 US-10-166-087-16	Sequence 16, Appl
15	38	45.2	35	15 US-10-092-750-1	Sequence 1, Appli

16	38	45.2	50	10 US-09-971-380-64	Sequence 64, Appl
17	38	45.2	64	10 US-09-971-380-62	Sequence 62, Appl
18	38	45.2	138	15 US-10-032-750-241	Sequence 241, App
19	38	45.2	327	15 US-10-102-806-570	Sequence 570, App
20	38	45.2	385	15 US-10-146-772-122	Sequence 122, App
21	38	45.2	571	9 US-09-815-242-11813	Sequence 11813, A
22	38	45.2	1053	9 US-09-841-132-583	Sequence 583, App
23	38	45.2	1053	14 US-10-007-693-97	Sequence 97, Appl
24	38	45.2	4840	15 US-10-156-761-10435	Sequence 10435, A
25	37.5	44.6	1265	15 US-10-198-070-69	Sequence 69, Appl
26	37	44.0	146	15 US-10-121-757B-12	Sequence 12, Appl
27	37	44.0	198	10 US-09-738-626-4812	Sequence 4812, Ap
28	37	44.0	334	10 US-09-794-715A-8	Sequence 8, Appli
29	37	44.0	334	12 US-10-286-581-8	Sequence 8, Appli
30	37	44.0	334	15 US-10-046-924-8	Sequence 8, Appli
31	37	44.0	426	9 US-09-815-242-5704	Sequence 5704, Ap
32	37	44.0	485	14 US-10-047-412A-4	Sequence 4, Appli
33	37	44.0	495	15 US-10-156-761-10045	Sequence 10045, A
34	37	44.0	515	15 US-10-106-698-4658	Sequence 4658, Ap
35	37	44.0	556	12 US-09-949-029-36	Sequence 36, Appl
36	37	44.0	575	9 US-09-839-136-2	Sequence 2, Appli
37	37	44.0	575	9 US-09-839-136-10	Sequence 10, Appl
38	37	44.0	705	9 US-09-815-242-12463	Sequence 12463, A
39	36.5	43.5	817	10 US-09-992-481-4	Sequence 4, Appli
40	36	42.9	61	9 US-09-925-301-1577	Sequence 1577, Ap
41	36	42.9	78	15 US-10-083-357-1237	Sequence 1237, Ap
42	36	42.9	80	15 US-10-106-698-5539	Sequence 5539, Ap
43	36	42.9	87	9 US-09-925-301-1433	Sequence 1433, Ap
44	36	42.9	95	10 US-09-925-300-1788	Sequence 1788, Ap
45	36	42.9	113	10 US-09-925-300-1789	Sequence 1789, Ap

ALIGNMENTS

RESULT 1

US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match 96.4% Score 81; DB 9; Length 204;
Best Local Similarity 93.8%; Pred. No. 1.4e-05;
Matches 15; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRRMSDEFEQ 16

Db 145 QRYGRLRRMTDEFEQ 160

RESULT 2

US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE


```
; Sequence 7, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/410,372
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 7:
US-09-894-657-7

Query Match      86.9%; Score 73; DB 9; Length 168;
Best Local Similarity 100.0%; Pred. No. 0.00023;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEF 14
      |||||
Db      108 QRYGRLRMSDEF 121

RESULT 7
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168

Query Match      86.9%; Score 73; DB 9; Length 168;
Best Local Similarity 100.0%; Pred. No. 0.00023;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEF 14
      |||||
Db      108 QRYGRLRMSDEF 121

RESULT 7
US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168

Query Match      86.9%; Score 73; DB 14; Length 168;
Best Local Similarity 100.0%; Pred. No. 0.00023;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRLRMSDEF 14
      |||||
Db      108 QRYGRLRMSDEF 121

RESULT 8
US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US20030068652A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECIFIC
; CONTEXT-INDEPENDENT ANTIBODIES COUPLED WITH DATABASE SEARCHIN
; FILE REFERENCE: CST-138 CIP3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147

Query Match      79.8%; Score 67; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.00017;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      4 GRELRMSDEFE 16
      |||||
Db      1 GRELRMSDEFE 13

RESULT 9
US-10-156-761-9145
; Sequence 9145, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
```

; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 9145
; LENGTH: 215
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-9145

Query Match 59.5%; Score 50; DB 15; Length 215;
Best Local Similarity 56.3%; Pred. No. 1.6;
Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDFEG 16
Db 108 ERWGGDLRRMRDEADG 123

RESULT 10

US-09-843-846-18
; Sequence 18, Application US/09843846
; Patent No. US20020042362A1
; GENERAL INFORMATION:
; APPLICANT: KUNSCH, CHARLES A
; CHOPRA, ARVIND
; ROSEN, CRAIG A
; TITLE OF INVENTION: HUMAN B-CELL TRANSLOCATION GENES-2 AND 3
; ANTIBODIES
; NUMBER OF SEQUENCES: 18
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX, P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, NW, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/843,846
; FILING DATE: 30-Apr-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/221,844
; FILING DATE: 29-DEC-1998
; APPLICATION NUMBER: US 08/718,738
; FILING DATE: 18-SEP-1996
; APPLICATION NUMBER: US 08/463,382
; FILING DATE: 05-JUN-1995
; APPLICATION NUMBER: US 08/460,104
; FILING DATE: 02-JUN-1995
; APPLICATION NUMBER: PCT/US95/03323
; FILING DATE: 17-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: STEFFE, ERIC K.
; REGISTRATION NUMBER: 36,688
; REFERENCE/DOCKET NUMBER: 1488.0730005/EKS/PSC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 213 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; SEQUENCE DESCRIPTION: SEQ ID NO: 18:

US-09-843-846-18
Query Match 51.2%; Score 43; DB 9; Length 213;

Best Local Similarity 43.8%; Pred. No. 21;
Matches 7; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDFEG 16
Db 22 RRFGEELERLKKYEG 37

RESULT 11

US-09-331-631A-22
; Sequence 22, Application US/09331631A
; Patent No. US20020168392A1
; GENERAL INFORMATION:
; APPLICANT: Manners, John M.
; APPLICANT: Marcus, John Paul
; APPLICANT: Goulter, Kenneth C.
; APPLICANT: Green, Jodie L.
; TITLE OF INVENTION: ANTIMICROBIAL PROTEINS
; FILE REFERENCE: CULLN23.001APC
; CURRENT APPLICATION NUMBER: US/09/331,631A
; CURRENT FILING DATE: 1999-06-21
; PRIOR APPLICATION NUMBER: PCT/AU97/00874
; PRIOR FILING DATE: 1997-12-22
; PRIOR APPLICATION NUMBER: AU PO 4275
; PRIOR FILING DATE: 1996-12-20
; NUMBER OF SEQ ID NOS: 40
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22
; LENGTH: 582
; TYPE: PRT
; ORGANISM: Maize
US-09-331-631A-22

Query Match 48.8%; Score 41; DB 10; Length 582;
Best Local Similarity 50.0%; Pred. No. 1.3e+02;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDFEG 16
Db 533 ERHGREEREKEERE 548

RESULT 12

US-10-156-761-13594
; Sequence 13594, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 13594
; LENGTH: 380
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-13594

Query Match 47.6%; Score 40; DB 15; Length 380;
Best Local Similarity 63.6%; Pred. No. 1.2e+02;
Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

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QY      3 YGRELRLMSDE 13
      | :|||:|:|
Db      241 YEKELRLADE 251

RESULT 13
US-10-146-473-41
; Sequence 41, Application US/10146473
; Publication No. US20030108888A1
; GENERAL INFORMATION:
; APPLICANT: Scanlan, Matthew
; APPLICANT: Gout, Ivan
; APPLICANT: Stockert, Elisabeth
; APPLICANT: Gure, Ali
; APPLICANT: Chen, Yao-Tseng
; APPLICANT: Old, Lloyd
; TITLE OF INVENTION: Breast Cancer Antigens
; FILE REFERENCE: L00461/70130(JRV)
; CURRENT APPLICATION NUMBER: US/10/146,473
; CURRENT FILING DATE: 2002-05-15
; PRIOR APPLICATION NUMBER: US 60/291,150
; PRIOR FILING DATE: 2001-05-15
; NUMBER OF SEQ ID NOS: 82
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 41
; LENGTH: 2871
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-146-473-41

Query Match      47.6%; Score 40; DB 15; Length 2871;
Best Local Similarity 60.0%; Pred. No. 9.7e+02;
Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY      1 QRYGRELRLMSDEFE 15
      || |||:|:|
Db      1648 QRTQELRLRLSSEVE 1662

RESULT 14
US-10-166-087-16
; Sequence 16, Application US/10166087
; Publication No. US2003007767A1
; GENERAL INFORMATION:
; APPLICANT: Ecopia Biosciences Inc.
; APPLICANT: Farnet, Chris
; APPLICANT: Staffa, Alfredo
; APPLICANT: Zazopoulos, Emmanuel
; TITLE OF INVENTION: Genes and proteins for the biosynthesis of anthramycin
; FILE REFERENCE: 3014-2US
; CURRENT APPLICATION NUMBER: US/10/166,087
; CURRENT FILING DATE: 2002-06-11
; NUMBER OF SEQ ID NOS: 51
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 16
; LENGTH: 764
; TYPE: PRT
; ORGANISM: Streptomyces refuineus subspecies thermotolerans
US-10-166-087-16

Query Match      45.8%; Score 38.5; DB 15; Length 764;
Best Local Similarity 62.5%; Pred. No. 4.3e+02;
Matches 10; Conservative 2; Mismatches 3; Indels 1; Gaps 1;

QY      1 QRYGRELRLMSDEFE 16
      || |||:|:|
Db      220 QRPGRRL-TYSNEYEG 234

RESULT 15
US-10-092-750-1
; Sequence 1, Application US/10092750
; Publication No. US20030032157A1

```

```

; GENERAL INFORMATION:
; APPLICANT: Hammond, Philip W.
; APPLICANT: Alpin, Julia
; APPLICANT: Wright, Martin C.
; TITLE OF INVENTION: Polypeptides Interactive with BCL-XL
; FILE REFERENCE: 50036/050002
; CURRENT APPLICATION NUMBER: US/10/092,750
; CURRENT FILING DATE: 2002-03-07
; PRIOR APPLICATION NUMBER: US 60/274,526
; PRIOR FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 253
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 35
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-092-750-1

Query Match      45.2%; Score 38; DB 15; Length 35;
Best Local Similarity 70.0%; Pred. No. 21;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY      5 RELRLMSDEF 14
      :|||:|:|
Db      19 QELRLIGDEF 28

Search completed: September 15, 2003, 17:47:52
Job time : 12.6857 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:18:16 ; Search time 7.2 Seconds
(without alignments)
213.708 Million cell updates/sec

Title: US-09-544-664-28
Perfect score: 84
Sequence: 1 QRYGRELRLMSDEFEQ 16

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_76.*

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Score	Match	Length	ID	Description
1	84	100.0	204	2 A55671	bad protein - mous
2	44	52.4	564	2 H75403	glycosyl hydrolase
3	43	51.2	463	2 H83036	probable two-compo
4	42.5	50.6	1014	2 T36031	excinuclease ABC c
5	42	50.0	370	2 S38185	2-dehydro-3-deoxy-
6	42	50.0	445	2 G97123	probable Fe-S oxid
7	41	48.8	84	2 F84388	hypothetical prote
8	41	48.8	113	2 JQ1128	gas-vesicle operon
9	41	48.8	113	2 T08234	gas-vesicle operon
10	41	48.8	220	2 F72289	oxidoreductase, so
11	41	48.8	271	2 AB1762	phosphate ABC tran
12	41	48.8	271	2 AH1386	phosphate ABC tran
13	41	48.8	322	1 RG3YC3	regulatory protein
14	41	48.8	398	2 G69496	ATP-dependent 26S
15	41	48.8	582	2 B53234	vicillin-like stora
16	40.5	48.2	334	2 A39172	Antho-Rfamidae neur
17	40.5	48.2	435	2 A44308	Antho-Rfamidae prec
18	40	47.6	206	2 C36365	transforming prote
19	40	47.6	219	2 T15466	hypothetical prote
20	40	47.6	374	2 C84338	spermidine/putresc
21	40	47.6	411	2 F87544	transcription regu
22	40	47.6	887	2 T38885	probable ATP-depen
23	40	47.6	967	2 F82668	oxoglutarate dehyd
24	40	47.6	2677	2 A38194	desmoplakin I - hu
25	40	47.6	5138	2 B96695	hypothetical prote
26	39.5	47.0	310	2 D86675	mevalonate kinase
27	39.5	47.0	910	2 G91024	NADH dehydrogenase
28	39.5	47.0	910	2 H85868	NADH dehydrogenase
29	39.5	47.0	910	2 A65000	NADH2 dehydrogenas

30 39.5 47.0 910 2 AI0796
31 39.5 47.0 914 2 AD0311
32 39 46.4 73 2 AC3365
33 39 46.4 87 2 A82928
34 39 46.4 91 2 T26996
35 39 46.4 105 2 B84184
36 39 46.4 232 2 A42095
37 39 46.4 275 2 C69808
38 39 46.4 315 2 T39444
39 39 46.4 331 2 E90121
40 39 46.4 365 2 S42107
41 39 46.4 380 2 T32163
42 39 46.4 501 2 H84727
43 39 46.4 574 2 T48113
44 39 46.4 1016 2 H75356
45 39 46.4 1248 2 G83278

ALIGNMENTS

RESULT 1

A55671
bad protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999
C:Accession: A55671
R:Yang, E.; Zha, J.; Jockel, J.; Boise, L.H.; Thompson, C.B.; Korsmeyer, S.J.
Cell 80, 285-291, 1995
A:Title: Bad, a heterodimeric partner for Bcl-X-L and Bcl-2, displaces Bax and promot
A:Reference number: A55671; MUID:95136361; PMID:7834748
A:Accession: A55671
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-204 <YAN>
A:Cross-references: GB:L37296; NID:g639778; PIDN:AAA64465.1; PID:g639779
C:Keywords: heterodimer

Query Match 100.0%; Score 84; DB 2; Length 204;
Best Local Similarity 100.0%; Pred. No. 1.9e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRLMSDEFEQ 16
|||||
Db 145 QRYGRELRLMSDEFEQ 160

RESULT 2

H75403
glycosyl hydrolase, family 13 - Deinococcus radiodurans (strain RL)
C:Species: Deinococcus radiodurans
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 17-Mar-2000
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, R.J.
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.;
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.
Science 286, 1571-1577, 1999
A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans RL.
A:Reference number: A75250; MUID:20036896; PMID:10567266
A:Accession: H75403
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-564 <WHI>
A:Cross-references: GB:AE001983; GB:AE000513; NID:g6459123; PIDN:AAF10944.1; PID:g645
C:Genetics:
A:Gene: DR1375
A:Map Position: 1
C:Superfamily: alpha-glucosidase; alpha-amylase core homology

Query Match 52.4%; Score 44; DB 2; Length 564;
Best Local Similarity 61.5%; Pred. No. 23;
Matches 8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 YGRLRMSDFE 15
| ||||| :|||
Db 283 YVREMRVYDFED 295

RESULT 3
H83036
probable two-component sensor PA4886 [Imported] - Pseudomonas aeruginosa (strain PA01)
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 14-Sep-2001
C:Accession: H83036
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Miziochichi, S.D.; Warrenner, P.; Hickey, M.J.; Boman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim, N.; Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic pathogen
A:Reference number: A82950; MUID:20437337; PMID:10984043
A:Accession: H83036
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-463 <STO>
A:Cross-references: GB:AE004901; GB:AE004091; NID:g9951147; PIDN:AAG08271.1; GSPDB:GN0011
A:Experimental source: strain PA01
A:Genetics:
A:Gene: PA4886
C:Superfamily: hypothetical protein HLL707; sensor histidine kinase homology

Query Match 51.2%; Score 43; DB 2; Length 463;
Best Local Similarity 61.5%; Pred. No. 28;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 4 GRLRMSDFEG 16
| ||||| :|||
Db 321 GEELRQAEYFEG 333

RESULT 4
T36031
excinuclease ABC chain A SCC54.18c [similarity] - Streptomyces coelicolor
N:Contains: excision endonuclease ABC (EC 3.1.-.-) chain A
C:Species: Streptomyces coelicolor
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 02-Feb-2001
C:Accession: t36031
R:Seeger, K.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
submitted to the EMBL Data Library, March 1999
A:Reference number: Z21581
A:Accession: T36031
A>Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-1014 <SEE>
A:Cross-references: EMBL:AL035591; PIDN:CAB38148.1; GSPDB:GN000070; SCOEDB:SCC54.18c
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: uvrA; SCOEDB:SCC54.18c
C:Superfamily: excinuclease ABC chain A; ATP-binding cassette homology
C:Keywords: ATP; DNA binding; DNA repair; hydrolase; nucleotide binding; P-loop
F:32-39/Region: nucleotide-binding motif A (P-loop)
F:645-652/Region: nucleotide-binding motif A (P-loop)

Query Match 50.6%; Score 42.5; DB 2; Length 1014;
Best Local Similarity 66.7%; Pred. No. 75;
Matches 10; Conservative 1; Mismatches 3; Indels 1; Gaps 1;

QY 2 RYGRLRMSDFEG 16
| ||||| || :|||
Db 368 RYGR-RYTTAFEG 381

RESULT 5
S38185
2-dehydro-3-deoxy-phosphoheptonate aldolase (EC 4.1.2.15) ARO4 - yeast (Saccharomyces cerevisiae)
N:Alternate names: 3-deoxy-D-arabino-heptulosonate-7-phosphate synthase; DAP synthase;

C:Species: Saccharomyces cerevisiae
C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 03-Jun-2002
C:Accession: S38185; S46126; S46130; JN0322; B48651
R:Doignon, F.; Biteau, N.; Aigle, M.; Crouzet, M.
Yeast 9, 1131-1137, 1993
A:Title: The complete sequence of a 6794 bp segment located on the right arm of chrom
A:Reference number: S38185; MUID:94078675; PMID:8256522
A:Accession: S38185
A>Status: translation not shown
A:Molecule type: DNA
A:Residues: 1-370 <DOI>
A:Cross-references: GB:L20296; NID:g3111101; PIDN:AAA65607.1; PID:g311102
R:Aljinovic, G.; Pohl, F.M.; Pohl, F.M.
submitted to the Protein Sequence Database, August 1994
A:Reference number: S45906
A:Accession: S46126
A:Molecule type: DNA
A:Residues: 1-370 <ALJ>
A:Cross-references: EMBL:Z36118; NID:g536664; PIDN:CAA85212.1; PID:g536665; MIPS:YBR2
R:Aigle, M.; Bactlet, M.C.; Barthe, C.; Biteau, N.; Crouzet, M.; Doignon, F.
submitted to the Protein Sequence Database, August 1994
A:Reference number: S45940
A:Accession: S46130
A:Molecule type: DNA
A:Residues: 1-370 <AIG>
A:Cross-references: EMBL:Z36118; NID:g536664; PIDN:CAA85212.1; PID:g536665; MIPS:YBR2
R:Kuenzler, M.; Paravicini, G.; Egli, C.M.; Paravicini, G.; Braus, G.H.
Gene 113, 67-74, 1992
A:Title: Cloning, primary structure and regulation of the ARO4 gene, encoding the tyr
A:Reference number: JN0322; MUID:92225349; PMID:1348717
A:Accession: JN0322
A:Molecule type: DNA
A:Residues: 1-204,208-370 <KUE>
A:Cross-references: EMBL:X61107
R:Kuenzler, M.; Balmelli, T.; Egli, C.M.; Paravicini, G.; Braus, G.H.
J. Bacteriol. 175, 5548-5558, 1993
A:Title: Cloning, primary structure, and regulation of the HIS7 gene encoding a bifun
A:Reference number: A48651; MUID:93374850; PMID:8366040
A:Accession: B48651
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 352-370 <KU2>
A:Cross-references: GB:X61107
C:Comment: This enzyme catalyzes the condensation of phosphoenolpyruvate and D-erythr
C:Genetics:
A:Gene: SGD:ARO4
A:Cross-references: SGD:S0000453; MIPS:YBR249c
A:Map position: 2R
C:Function:
A:Description: aldehyde-lyase; carbon-carbon lyase
A:Pathway: aromatic amino acid biosynthesis; shikimate pathway
A:Note: first step in shikimate pathway
C:Superfamily: phospho-2-dehydro-3-deoxyheptonate aldolase
C:Keywords: aldehyde-lyase; aromatic amino acid biosynthesis; carbon-carbon lyase; cy

Query Match 50.0%; Score 42; DB 2; Length 370;
Best Local Similarity 43.8%; Pred. No. 32;
Matches 7; Conservative 4; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDFEG 16
| | | :||| :|||
Db 85 QEYALRLKSLDELKG 100

RESULT 6
G97123
Probable Fe-S oxidoreductase CAC1813 [Imported] - Clostridium acetobutylicum
C:Species: Clostridium acetobutylicum
C:Date: 14-Sep-2001 #sequence_revision 14-Sep-2001 #text_change 30-Sep-2001
C:Accession: G97123
R:Nolling, J.; Breton, G.; Omelchenko, M.V.; Markarova, K.S.; Zeng, Q.; Gibson, R.; L
.; Daly, M.J.; Bennett, G.N.; Koonin, E.V.; Smith, D.R.
J. Bacteriol. 183, 4823-4836, 2001

Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Nature
Globe

TOW

A;Authors: Utterback, T.; Cotton, M.D.; Spriggs, T.; Artiach, P.; Kaine, B.P.; Sykes, S.
Smith, H.O.; Woese, C.R.; Venter, J.C.
A;Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archaeon
A;Reference number: A69250; MUID:98049343; PMID:9389475
A;Accession: G69496
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-398 <KLE>
A;Cross-references: GB:AE000967; GB:AE000782; NID:g2689290; PIDN:AA89280.1; PID:g264858
C;Superfamily: ATP-dependent 36S proteinase; FtsH/SEC18/CDC48-type ATP-binding domain ho
C;Keywords: ATP; nucleotide binding; P-loop
F:155-365/Domain: FtsH/SEC18/CDC48-type ATP-binding domain homology <VATP>
F:182-189/Region: nucleotide-binding motif A (P-loop)

Query Match 48.8%; Score 41; DB 2; Length 398;
Best Local Similarity 57.1%; Pred. NO. 51;
Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 RYGRRLRMSDEPE 15
Db 43 RYREVRRLRSEVE 56
|||:|:|:|

RESULT 15

B53234
vicilin-like storage protein Gbl-L, embryo - maize
N;Alternate names: globulin-1L
C;Species: Zea mays (maize)
C;Date: 02-May-1994 #sequence_revision 18-Nov-1994 #text_change 24-Nov-1999
C;Accession: B53234; S21824
R;Belanger, F.C.; Kriz, A.L.
Genetics 129, 863-872, 1991
A;Title: Molecular basis for allelic polymorphism of the maize Globulin-1 gene.
A;Reference number: A53234; MUID:92090707; PMID:1752424
A;Accession: B53234
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-582 <BEL>
A;Cross-references: EMBL:X59083; NID:g22283; PIDN:CAA1809.1; PID:g22284
A;Experimental source: Inbred line W64A6
A;Note: sequence extracted from NCBI backbone (NCBIP:71285)
C;Genetics:
A;Gene: Gbl-L
A;Introns: 167/1; 225/3; 252/3; 349/3
C;Superfamily: glycinin

Query Match 48.8%; Score 41; DB 2; Length 582;
Best Local Similarity 50.0%; Pred. NO. 75;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRRLRMSDEFE 16
Db 533 ERHGREEREKEFE 548
:|:|:|:| :|:|

Search completed: September 15, 2003, 17:27:01
Job time : 7.2 secs

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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:55 ; Search time 3.77143 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-28
Perfect score: 84
Sequence: 1 QRYGRELRRMSDEFEG 16

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	84	100.0	204	1 BAD_MOUSE	Q61337 mus musculus
2	84	100.0	205	1 BAD_RAT	O35147 rattus norv
3	73	86.9	168	1 BAD_HUMAN	Q92934 homo sapien
4	42.5	50.6	1014	1 UVR_A_STRCO	O92507 streptomyce
5	42	50.0	196	1 BIM_MOUSE	O54918 mus musculus
6	42	50.0	196	1 BIM_RAT	O84998 rattus norv
7	42	50.0	370	1 AROG_YEAST	P32449 saccharomyc
8	42	50.0	653	1 HT2A_HUMAN	Q13049 homo sapien
9	41	48.8	113	1 GVK1_HALNL	P24375 halobacteri
10	41	48.8	220	1 6PGL_THEMA	Q9x08 thermotoga
11	41	48.8	322	1 SNF4_YEAST	P12904 saccharomyc
12	41	48.8	328	1 SNF4_KLJLA	Q9P869 kluyveromyc
13	41	48.8	398	1 PSME_ARCFU	O28303 archaeoglob
14	40.5	48.2	334	1 FMRA_CALPA	Q01133 callitactis
15	40.5	48.2	429	1 FMR2_ANTEL	Q16994 anthopleura
16	40.5	48.2	435	1 FMRL_ANTEL	P10419 anthopleura
17	40	47.6	205	1 RAS3_RHRA	P22280 rhizomucor
18	40	47.6	887	1 YAY5_SCHPO	Q10213 schizosacch
19	40	47.6	2871	1 DESP_HUMAN	P15924 homo sapien
20	39.5	47.0	907	1 NUOG_ECOLI	P33602 escherichia
21	39.5	47.0	907	1 NUOG_SALTY	P33900 salmonella
22	39	46.4	87	1 Y152_UREPA	Q9P322 ureaplasma
23	39	46.4	95	1 R37A_HALNL	O9B968 halobacteri
24	39	46.4	232	1 AP3_ARATH	P35832 arabidopsis
25	39	46.4	365	1 RAS1_SCHPO	P36601 schizosacch
26	39	46.4	889	1 SEC3_DROME	Q9VW94 drosophila
27	39	46.4	1016	1 UVR_A_DEIRA	Q46577 deinococcus
28	39	46.4	1535	1 LML1_CABEL	O18823 caenorhabdi
29	39	46.4	8797	1 SNE1_HUMAN	O8nf91 homo sapien
30	38.5	45.8	323	1 FRG1_CABEL	O18282 caenorhabdi
31	38.5	45.8	468	1 SELA_PSEAE	Q9H01 pseudomonas
32	38	45.2	177	1 IF3_CLOPE	O8XJ67 clostridium
33	38	45.2	198	1 BIM_HUMAN	O43521 homo sapien

RESULT 1

BAD_MOUSE ID	BAD_MOUSE STANDARD;	PRT;	204 AA.
AC Q61337;			
DT 01-NOV-1997 (Rel. 35, Created)			
DT 01-NOV-1997 (Rel. 35, Last sequence update)			
DT 28-FEB-2003 (Rel. 41, Last annotation update)			
DE Bcl-2 antagonist of cell death (BAD) (Bcl-2 binding component			
DE 6) (Bcl-XL/Bcl-2 associated death promoter).			
GN BAD OR BOC6.			
OS Mus musculus (Mouse).			
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.			
OX NCBI_TaxID=10090;			
RN [1]			
RP SEQUENCE FROM N.A.			
RC TISSUE=Brain, and Thymus;			
RX MEDLINE=95136361; PubMed=7834748;			
RA Yang E., Zha J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;			
RT "Bad, a heterodimeric partner for Bcl-XL and Bcl-2, displaces Bax and			
RT promotes cell death.";			
RL Cell 80:285-291(1995).			
RN [2]			
RP PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.			
RX MEDLINE=9802383; PubMed=9381178;			
RA Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;			
RT "Interleukin-3-induced phosphorylation of BAD through the protein			
RT kinase Akt.";			
RL Science 278:687-689(1997).			
RN [3]			
RP MUTAGENESIS OF SERINE RESIDUES.			
RX MEDLINE=20403302; PubMed=10949026;			
RA Datta S.R., Katsov A., Hu L., Petros A., Fesik S.W., Yaffe M.B.,			
RA Greenberg M.E.;			
RT "14-3-3 proteins and survival kinases cooperate to inactivate BAD by			
RT BH3 domain phosphorylation.";			
RL Mol. Cell 6:41-51(2000).			
CC -1- FUNCTION: Promotes cell death. Successfully competes for the			
CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level			
CC of heterodimerization of these proteins with BAX. Can reverse the			
CC death repressor activity of Bcl-x(L), but not that of Bcl-2.			
CC Appears to act as a link between growth factor receptor signaling			
CC and the apoptotic pathways.			
CC -1- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-			
CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).			
CC The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.			
CC -1- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon			
CC phosphorylation, locates to the cytoplasm.			
CC -1- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND			
CC BAX for their pro-apoptotic activity and for their interaction			
CC with anti-apoptotic members of the Bcl-2 family.			
CC -1- PTM: Phosphorylated on Ser-112 in response to survival stimuli.			
CC Subsequent phosphorylation on Ser-136 promotes heterodimerization			
CC with 14-3-3 proteins. This interaction then facilitates the			
CC phosphorylation at Ser-155, a site within the BH3 domain, leading			
CC to the release of Bcl-x(L) and the promotion of cell survival.			

34	38	45.2	207	1	THIE_PYRAB	Q9uzg5 pyrococcus
35	38	45.2	370	1	AROG_CANAL	P79023 candida alb
36	38	45.2	570	1	STIM_DROME	P83094 drosophila
37	38	45.2	712	1	PSEL_HUMAN	P22079 homo sapien
38	38	45.2	861	1	GER3_YEAST	P34160 saccharomyc
39	38	45.2	1047	1	RIR1_CHLMO	Q9P193 chlamydia m
40	38	45.2	1268	1	RIR1_CHLTR	O84834 chlamydia t
41	38	45.2	1268	1	VGLN_HUMAN	Q00341 homo sapien
42	38	45.2	1270	1	VGLN_CHICK	P81021 gallus gall
43	38	45.2	1958	1	UVR1_SCHPO	O60152 schizosacch
44	37.5	44.6	992	1	UVR1_MICLU	P33567 micrococcus
45	37	44.0	126	1	LEO3_BUCUL	Q9aqc8 buchnera ap

ALIGNMENTS

Query Match 100.0%; Score 84; DB 1; Length 205;
 Best Local Similarity 100.0%; Pred. No. 1.3e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRRMSDFEG 16
 |||||
 DB 146 QRYGRLRRMSDFEG 161

RESULT 3
 ID BAD_HUMAN STANDARD; PRT; 168 AA.
 AC Q2934; O14803;
 DT 01-NOV-1997 (rel. 35, Created)
 DT 16-OCT-2001 (rel. 40, Last sequence update)
 DT 15-SEP-2003 (rel. 42, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-2
 DE XI/Bcl-2 associated death promoter) (BCL2-like 8 protein).
 GN BAD OR BBC6 OR BCL2L8.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 CX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Yin D.X., Li Z., Huang B., Chen S., Zhou H.;
 RT "A human protein that interacts with Bcl-2 and have homology to mouse
 RT BAD";
 RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.
 RX MEDLINE=97083574; PubMed=8929532;
 RA Wang H.-G., Rapp U.R., Reed J.C.;
 RT "Bcl-2 targets the protein kinase Raf-1 to mitochondria";
 RL Cell 87:629-638(1996).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Takayama S., Reed J.C.;
 RL Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A., AND DIMERIZATION.
 RX TISSUE=Bone marrow;
 RC MEDLINE=98049554; PubMed=9388232;
 RA Orlille S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,
 RA Chang S., Weeks S., Fritz L.C., Oltersdorf T.;
 RT "Dimerization properties of human BAD";
 RL J. Biol. Chem. 272:30866-30872(1997).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lung;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg K.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Udén T.B., Toshlyuk S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalls D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [6]

RP STRUCTURE BY NMR OF 103-127.
 RX MEDLINE=21073561; PubMed=11206074;
 RA Petros A.M., Nettekheim D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
 RA Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
 RA Pesik S.W.;
 RT "Rationale for Bcl-xL/Bad peptide complex formation from structure,
 RT mutagenesis, and biophysical studies";
 RL Protein Sci. 9:2528-2534(2000).
 CC -I- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 CC similarity). Appears to act as a link between growth factor
 CC receptor signaling and the apoptotic pathways.
 CC -I- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 CC The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
 CC similarity).
 CC -I- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm.
 CC -I- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
 CC -I- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -I- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-99 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-118, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-99 is the major site of AKT/PKB phosphorylation, Ser-118 the
 CC major site of protein kinase A (CAK) phosphorylation (by
 CC similarity).
 CC -I- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
 CC -I- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -I- CAUTION: Ref.1 sequence differs from that shown due to frameshifts
 CC in position 64 and 91.
 CC -----
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 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; U66879; AAB36516.1; ALT_FRAME.
 CC EMBL; AF021792; AAB72092.1; -
 CC EMBL; AF031523; AAB88124.1; -
 CC EMBL; BC001901; AAB01901.1; -
 CC PDB; 1G5J; 07-FEB-01.
 CC Genew; HGNC:936; BAD.
 CC MIM; 603167; -
 CC GO; GO:0005737; Cytoplasm; NAS.
 CC GO; GO:0005741; C-mitochondrial outer membrane; NAS.
 CC GO; GO:0005515; F-protein binding activity; NAS.
 CC GO; GO:0008632; P-apoptotic program; TAS.
 CC GO; GO:0006917; P-induction of apoptosis; NAS.
 CC InterPro; IPR000712; Bcl2_BH.
 CC PROSITE; PS01259; BH3; FALSE_NEG.
 CC Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
 KW DOMAIN 110 124
 FT MOD_RES 75 75
 FT PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).
 FT MOD_RES 99 99
 FT PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).
 FT MOD_RES 118 118
 FT PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).
 FT VARIANT 107 107
 FT A -> S (in dbSNP:3729933).
 FT /FTID=VAR_015380.
 FT HELIX 106 121
 FT SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
 SQ
 Query Match 86.9%; Score 73; DB 1; Length 168;

Best Local Similarity 100.0%; Pred. No. 6.7e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDEF 14
|||||
Db 108 QRYGRLRMSDEF 121

RESULT 4
ID UVR_A_STRCO STANDARD; PRT; 1014 AA.
AC Q92507;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE UVRABC system protein A (UvrA protein) (Excinuclease ABC subunit A).
GN UVR A OR SCO1958 OR SCC54.18C.
OS Streptomyces coelicolor.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=1902;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=A3(2) / M145;
RC MEDLINE=21996410; PubMed=12000953;
RA Bentley S.D., Chater K.F., Harris D.E., Quail M.A., Kieser H.,
RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Collins M.,
RA Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,
RA Cronin A., Fraser A., Goble A., Hidalgo J., Hornsby T., Howarth S.,
RA Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neill S.,
RA Rabinowitch E., Rajandream M.A., Rutherford K., Rutter S.,
RA Seeger K., Saunders D., Sharp S., Squares R., Taylor K.,
RA Warren T., Wietzorek A., Woodward J., Bartell B.G., Parkhill J.,
RA Hopwood D.A.;
RI "Complete genome sequence of the model actinomycete Streptomyces
coelicolor A3(2).";
RL Nature 417:141-147(2002).
CC -!- FUNCTION: The UvrABC repair system catalyzes the recognition and
processing of DNA lesions. UvrA is an ATPase and a DNA-binding
protein. A damage recognition complex composed of 2 uvrA and 2
uvrB subunits scans DNA for abnormalities. When the presence of a
lesion has been verified by uvrB, the uvrA molecules dissociate
(by similarity).
CC -!- SUBUNIT: Forms a heterotetramer with uvrB during the search for
lesions (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: BELONGS TO THE ABC TRANSPORTER FAMILY. UVR A SUBFAMILY.
CC
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or send an email to license@isb-sib.ch).
CC
CC EMBL; AL035591; CAB38148.1; -;
CC PIR; T36031; T36031.
CC HAMAP; MF_00205; -; 1.
CC InterPro; IPR003439; ABC_transporter.
CC InterPro; IPR004602; UvrA.
CC Pfam; PF00005; ABC_tran; 2.
CC ProDom; PD00006; ABC_transporter; 1.
CC TIGRFAMs; TIGR00630; uvrA; 1.
CC PROSITE; PS00211; ABC_TRANSPORTER_1; 2.
CC PROSITE; PS08933; ABC_TRANSPORTER_2; 1.
CC SOS response; Exclusion nuclease; DNA repair; DNA recombination;
KW DNA excision; ATP-binding; DNA-binding; Repeat; zinc; Metal-binding;
KW Zinc-finger; Complete proteome.
FT NP_BIND 32 39 ATP (POTENTIAL).
FT NP_BIND 645 652 ATP (POTENTIAL).
FT ZN_FING 744 770 C4-TYPE.
FT SEQUENCE 1014 AA; 110997 MW; 084D6B18692A792D CRC64;

Query Match 50.8%; Score 42.5; DB 1; Length 1014;
Best Local Similarity 66.7%; Pred. No. 39;
Matches 10; Conservative 1; Mismatches 3; Indels 1; Gaps 1;

QY 2 RYGRELRMSDEFG 16
|||||
Db 368 RYGRE-RYTTAFEG 381

RESULT 5
ID BIM_MOUSE STANDARD; PRT; 196 AA.
AC O54918; O54919; O54920;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE BCL2-like protein 11 (BCL2 interacting mediator of cell death).
GN BCL2L11 OR BIM.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.; FUNCTION. SUBCELLULAR LOCATION, TISSUE
SPECIFICITY, AND ALTERNATIVE SPLICING.
RX MEDLINE=98094360; PubMed=9430630;
RA O'Connor L., Strasser A., O'Reilly L.A., Hausmann G., Adams J.M.,
RA Cory S., Huang D.C.S.;
RL "Bim: a novel member of the Bcl-2 family that promotes apoptosis.";
EMBO J. 17:384-395(1998).
CC -!- FUNCTION: INDUCES APOPTOSIS. THE ISOFORMS VARY IN CYTOTOXICITY
WITH ISOFORM BIM BEING THE MOST POTENT AND ISOFORM BIMEL BEING
THE LEAST POTENT.
CC -!- SUBUNIT: FORMS HETERODIMERS WITH A NUMBER OF ANTIAPOPTOTIC BCL-2
PROTEINS INCLUDING MCL-1, BCL-2, BCL-XL, BFL-1, AND BHRF-1. DOES
NOT HETERODIMERIZE WITH PROAPOPTOTIC PROTEINS SUCH AS BAD, BOK,
BAX OR BAK (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: ASSOCIATED WITH INTRACYTOSOLIC MEMBRANES.
CC -!- ALTERNATIVE PRODUCTS:
Event-Alternative splicing; Named isoforms=3;
Name=BimEL;
IsoId=O54918-1; Sequence=Displayed;
Name=BimL;
IsoId=O54918-2; Sequence=VSP_000536;
Name=BimS;
IsoId=O54918-3; Sequence=VSP_000537;
CC -!- TISSUE SPECIFICITY: EXPRESSED IN A NUMBER OF B-AND T-LYMPHOID CELL
LINES.
CC -!- DOMAIN: THE BH3 DOMAIN IS REQUIRED FOR BCL-2 BINDING AND
CYTOTOXICITY.
CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
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CC
CC EMBL; AF032459; AAC40029.1; -;
CC EMBL; AF032460; AAC40030.1; -;
CC EMBL; AF032461; AAC40031.1; -;
CC MGD; MGI:1197519; Bcl2l11.
CC InterPro; IPR000712; Bcl2_BH.
KW Apoptosis; PS01259; BH3; FALSE_NEG.
FT DOMAIN 146 160 BH3.
FT VARSPPLIC 42 97 Missing (in isoform BimL).
FT VARSPPLIC 42 127 /FTId=VSP_000536.
FT /FTId=VSP_000537.

SQ SEQUENCE 196 AA; 22066 MW; 531C17655FIAC9AA CRC64;

Query Match 50.0%; Score 42; DB 1; Length 196;
Best Local Similarity 61.5%; Pred. No. 8.6;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 2 RYGRRLRMSDEF 14
DB 145 RIAQLRRIGDEF 157

RESULT 6

BIM_RAT
ID BIM_RAT STANDARD; PRT; 196 AA.
AC O88498; O88497; Q9WU18;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE BCL2-like protein 11 (BCL2 interacting mediator of cell death)
DE (Bcl-2 related ovarian death protein).
GN BCL2L1 OR BIM OR BOD.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RN [2]
RP SEQUENCE FROM N.A., FUNCTION, SUBUNIT, AND TISSUE SPECIFICITY
RP (ISOFORMS BOD-L; BOD-M AND BOD-S).
RC TISSUE-Ovary;
RX MEDLINE=98400436; PubMed=9731710;
RA Hsu S.Y., Lin P., Hsueh A.J.W.;
RT "BOD (Bcl-2-related ovarian death gene) is an ovarian BH3 domain-containing proapoptotic Bcl-2 protein capable of dimerization with diverse antiapoptotic Bcl-2 members.";
RL Mol. Endocrinol. 12:1437-1440(1998).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM BIML).
RA Chen D., Simon R.P., Chen J.;
RT "Cloning of rat bimL and bimL, and their differential expression in ischemia and normal rat brain.";
RL Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INDUCES APOPTOSIS.
CC -!- SUBUNIT: FORMS HETERODIMERS WITH A NUMBER OF ANTIAPOPTOTIC BCL-2 PROTEINS INCLUDING MCL-1, BCL-2, BCL-XL, BCL-L, AND BHRF-1. DOES NOT HETERODIMERIZE WITH PROAPOPTOTIC PROTEINS SUCH AS BAD, BOK, BAX OR BAK.
CC -!- SUBCELLULAR LOCATION: ASSOCIATED WITH INTRACYTOSOLASMIC MEMBRANES (BY SIMILARITY).
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=3;
CC Name=BOD-L;
CC IsoId=O88498-1; Sequence=Displayed;
CC Note=Isoform BOD-S is produced by alternative initiation at Met-104 of Isoform BOD-L;
CC Name=BimL;
CC IsoId=O88498-2; Sequence=VSP_000538;
CC Name=BOD-M;
CC IsoId=O88498-3; Sequence=VSP_000539;
CC Event=Alternative initiation;
CC Comment=2 isoforms, BOD-L (shown here) and BOD-S, are produced by alternative initiation at Met-1 and Met-104;
CC -!- TISSUE SPECIFICITY: Widely expressed.
CC -!- DOMAIN: THE BH3 DOMAIN IS REQUIRED FOR BCL-2 BINDING AND CYTOTOXICITY.
CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC
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CC EMBL; AF065433; AAC23595.1; -;
DR EMBL; AF065431; AAC23593.1; -;
DR EMBL; AF065432; AAC23594.1; -;
DR EMBL; AF136927; AAD26594.1; -;
DR InterPro; IPR000712; Bcl2.BH.
DR PROSITE; PS01259; BH3; FALSE.NEG.
KW Apoptosis; Alternative splicing; Membrane; Alternative initiation.
FT CHAIN 1 196
FT CHAIN 104 196
FT INIT_MET 104 104
FT DOMAIN 146 160
FT VARSPLIC 42 97
FT Missing (in isoform BimL).
FT VARSPLIC 42 127
FT Missing (in isoform BOD-M).
FT CONFLICT 136 136
FT E -> D (IN REF. 1; AAC23594).
SQ SEQUENCE 196 AA; 22055 MW; B4D2146F9C0B37A0 CRC64;

Query Match 50.0%; Score 42; DB 1; Length 196;
Best Local Similarity 61.5%; Pred. No. 8.6;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 2 RYGRRLRMSDEF 14
DB 145 RIAQLRRIGDEF 157

RESULT 7

AROG_YEAST
ID AROG_YEAST STANDARD; PRT; 370 AA.
AC P32449;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Phospho-2-dehydro-3-deoxyheptonate aldolase, tyrosine-inhibited (EC 4.1.2.15). (Phospho-2-keto-3-deoxyheptonate aldolase) (DAHP synthetase) (3-deoxy-D-arabino-heptulosonate 7-phosphate synthase).
GN ARO4 OR YBR249C OR YBR1701.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RN [2]
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=92225349; PubMed=1348717;
RA Kuenzler M., Paravicini G., Egli C., Irniger S., Braus G.H.;
RT "Cloning, primary structure and regulation of the ARO4 gene, encoding the tyrosine-inhibited 3-deoxy-D-arabino-heptulosonate-7-phosphate synthase from Saccharomyces cerevisiae.";
RL Gene 113:67-74(1992).
RN [2]
RP REVISIONS TO 205-207.
RA Kuenzler M.;
RN Submitted (NOV-1993); to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=S288C;
RX MEDLINE=94078675; PubMed=8256522;
RA Doignon F., Biteau N., Aigle M., Crouzet M.;
RT "The complete sequence of a 6794 bp segment located on the right arm of chromosome II of Saccharomyces cerevisiae. Finding of a putative tufPase in a yeast.";
RL Yeast 9:1131-1137(1993).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=S288C;
RA Aljinovic G., Pohl F.M., Pohl T.M.;
RN Submitted (AUG-1994) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: STEREOSPECIFIC CONDENSATION OF PHOSPHENOLPYRUVATE (PEP) AND D-ERYTHROSE-4-PHOSPHATE (E4P) GIVING RISE TO 3-DEOXY-D-ARABINO-HEPTULOSONATE-7-PHOSPHATE (DAHP).
CC -!- CATALYTIC ACTIVITY: 2-dehydro-3-deoxy-D-arabino-heptonate 7-


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CC phosphate + phosphate = phosphoenolpyruvate + D-erythrose 4-
CC phosphate + H(2)O.
CC -!- ENZYME REGULATION: INHIBITED BY TYROSINE.
CC -!- PATHWAY: Aromatic amino acids biosynthesis; shikimate pathway;
CC first step.
CC -!- INDUCTION: By amino acid starvation.
CC -!- SIMILARITY: BELONGS TO CLASS-I DHP SYNTHETASE FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: X61107; CAA43419.1; -.
DR GO: 0003849; F:2-dehydro-3-deoxyphosphoheptonate aldolase . . . ; IDA.
DR EMBL: L20296; AAA65607.1; -.
DR EMBL: X36118; CAA85212.1; -.
DR PIR: S38185; S38185.
DR HSSP: P00886; 1OR7.
DR SGD: S0000453; ARO4.
DR GO: 0003849; F:2-dehydro-3-deoxyphosphoheptonate aldolase . . . ; IDA.
DR InterPro: IPR006219; AroFGH.
DR InterPro: IPR006218; DAHP1/KDSA.
DR Pfam: PF00793; DAHP_synth_1; 1.
DR ProDom: PD005060; AroFGH; 1.
DR TIGRFAMs: TIGR00034; aroFGH; 1.
KW Aromatic amino acid biosynthesis; Lyase; Multigene family.
SQ SEQUENCE 370 AA; 39749 MW; 594ED48F24175979 CRC64;

Query Match 50.0%; Score 42; DB 1; Length 370;
Best Local Similarity 43.8%; Pred. No. 17;
Matches 7; Conservative 4; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRLRRMSDFEG 16
DQ 85 QYALRLKLKLSDELKG 100

RESULT 8
HT2A_HUMAN STANDARD; PRT; 653 AA.
AC Q13049; O9NQ8;
DT 01-NOV-1997 (Rel. 35, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Zinc-finger protein HT2A (72 kDa Tat-interacting protein) (Tripartite
DE motif-containing protein 32).
GN TRIM32 OR HT2A
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=95297135; PubMed=7778269;
RA Fridell R.A., Harding L.S., Bogerd H.P., Cullen B.R.;
RT Identification of a novel human zinc finger protein that
RT specifically interacts with the activation domain of lentiviral Tat
RT proteins.
RL Virology 209:347-357(1995).
RN [2]
RP SEQUENCE FROM N.A.
RA Sehra H.;
RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Skin;
RX Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,

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RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavini T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A.C., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield J.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- FUNCTION: MAY PLAY A SIGNIFICANT ROLE IN MEDIATING THE BIOLOGICAL
CC ACTIVITY OF THE HIV-1 TAT PROTEIN IN VIVO. BINDS SPECIFICALLY TO
CC THE ACTIVATION DOMAIN OF HIV-1 TAT AND CAN ALSO INTERACT WITH THE
CC HIV-2 AND EIAV TAT PROTEINS IN VIVO.
CC -!- SUBCELLULAR LOCATION: Nuclear.
CC -!- TISSUE SPECIFICITY: SPLEEN, THYMUS, PROSTATE, TESTIS, OVARY,
CC INTESTINE AND COLON.
CC -!- SIMILARITY: Contains 1 RING-type zinc finger.
CC -!- SIMILARITY: Contains 1 B box-type zinc finger.
CC -----
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CC -----
DR EMBL: U18543; AAA86474.1; -.
DR EMBL: AL133284; CAB92723.1; -.
DR EMBL: BC003154; AAH03154.1; -.
DR HSSP: P29590; 1BOR.
DR Genew: HGNC:16380; TRIM32.
DR MIM: 602290; -.
DR GO: 0005634; C:nucleus; TAS.
DR GO: 0003713; P:transcription co-activator activity; TAS.
DR InterPro: IPR001258; NHL.
DR InterPro: IPR000315; Znf_Box.
DR Pfam: PF01436; NHL; 5.
DR Pfam: PF00643; zf-B_box; 1.
DR Pfam: PF00097; zf-C3HC4; 1.
DR SMART: SM00336; BBOX; 1.
DR SMART: SM00184; RING; 1.
DR PROSITE: PS00119; ZF_BOX; 1.
DR PROSITE: PS00518; ZF_RING_1; 1.
DR PROSITE: PS00089; ZF_RING_2; 1.
KW Zinc-finger; Nuclear protein.
FT DOMAIN 2 POLY-ALA.
FT ZN_FING 20 65 RING-TYPE.
FT ZN_FING 103 133 B_BOX-TYPE.
FT CONFLICT 27 27 F -> I (IN REF. 1).
SQ SEQUENCE 653 AA; 71988 MW; D83B1595CA8378FD CRC64;

Query Match 50.0%; Score 42; DB 1; Length 653;
Best Local Similarity 61.5%; Pred. No. 30;
Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 QRYGRLRRMSDE 13
DQ 186 QYGHERRVQDE 198

RESULT 9
GVKL_HALN1
ID GVKL_HALN1 STANDARD; PRT; 113 AA.

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P24375; Q9H126;
 DT 01-MAR-1992 (Rel. 21, Created)
 DT 01-MAR-1992 (Rel. 21, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE GvpK protein 1.
 GN (GVPK11 OR GVPK OR VNG5021G) AND (GVPK12 OR VNG6021G).
 OS Halobacterium sp. (strain NRC-1 / ATCC 700922 / JCM 11081), and
 OS Halobacterium sp. (strain NRC-817).
 OG Plasmid pNRC100, Plasmid pNRC200, and plasmid pH1.
 OC Archaea; Euryarchaeota; Halobacteria; Halobacteriales;
 OC Halobacteriaceae; Halobacterium.
 OX NCBI_TaxID=64091, 148370;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NRC-1; PLASMID=pNRC100;
 RX MEDLINE=91323716; PubMed=1864501;
 RA Jones J.G., Young D.C., Dassarma S.;
 RT "Structure and organization of the gas vesicle gene cluster on the
 RT Halobacterium halobium plasmid pNRC100.";
 RL Gene 102:117-122(1991).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NRC-1; PLASMID=pNRC100;
 RX MEDLINE=99063795; PubMed=9847077;
 RA Ng W.V., Clufo S.A., Smith T.M., Bumgarner R.E., Baskin D., Faust J.,
 RA Hall B., Loretz C., Seto J., Slagel J., Hood L., DasSarma S.;
 RT "Snapshot of a large dynamic replicon in a halophilic archaeon:
 RT megaplasmid or minichromosome?";
 RL Genome Res. 8:1131-1141(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NRC-1; PLASMID=pNRC200;
 RX MEDLINE=20504483; PubMed=11016950;
 RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,
 RA Saikia H.D., Lasky S.R., Baliga N.S., Thorsson V., Sbrogna J.,
 RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
 RA Leitzhauser B., Keller K., Cruz R., Danson M.J., Hough D.W.,
 RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
 RA Isenbarger T.A., Peck K.F., Pohlschroder M., Spudis J.L., Jung K.-H.,
 RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
 RA Ehardt H., Lowe T.M., Liang P., Riley M., Hood L., DasSarma S.;
 RT "Genome sequence of Halobacterium species NRC-1.";
 RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NRC-817; PLASMID=pH1;
 RX MEDLINE=92065812; PubMed=1956294;
 RA Horne M., Englert C., Wimmer C., Pfeifer F.;
 RT "A DNA region of 9 kbp contains all genes necessary for gas vesicle
 RT synthesis in halophilic archaeobacteria.";
 RL Mol. Microbiol. 5:1159-1174(1991)
 CC -1- FUNCTION: MAY PLAY A STRUCTURAL OR REGULATORY ROLE IN GAS
 CC VESICLE SYNTHESIS.
 CC -----
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 CC -----
 CC DR EMBL; M58557; AAA98188.1; -;
 CC DR EMBL; AF016485; AAC82801.1; -;
 CC DR EMBL; AF005141; AAG20718.1; -;
 CC DR EMBL; X55648; CAA39178.1; -;
 CC DR PIR; T08234; T08234.
 CC DR Pfam; PF05121; GvpK; 1.
 CC KW Gas vesicle; Plasmid; Complete proteome.
 CC SQ SEQUENCE 113 AA; 12695 MW; 97A469D2C1643ABF CRC64;
 CC -----
 CC Query Match 48.8%; Score 41; DB 1; Length 113;
 CC Best Local Similarity 46.7%; Pred. No. 7;
 CC Matches 6; Conservative 6; Mismatches 3; Indels 0; Gaps 0;

Matches 7; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
 Qy 1 QRYGRELRRMSDEFE 15
 Db 50 ERLGRLQALEDELE 64
 RESULT 10
 6PGL_THEME
 ID 6PGL_THEME STANDARD; PRT; 220 AA.
 AC QXON8;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 6-phosphogluconolactonase (EC 3.1.1.31) (6PGL).
 GN PGL OR DEVB OR TML154.
 OS Thermotoga maritima.
 OC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
 OX NCBI_TaxID=2336;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MSB8 / DSM 3109;
 RX MEDLINE=99287316; PubMed=10360571;
 RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
 RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
 RA McDonald L., Utterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
 RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
 RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
 RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
 RT "Evidence for lateral gene transfer between Archaea and Bacteria from
 RT genome sequence of Thermotoga maritima.";
 RL Nature 399:323-329(1999).
 CC -1- FUNCTION: HYDROLYSIS OF 6-PHOSPHOGLUCONOLACTONE TO 6-
 CC PHOSPHOGLUCONATE.
 CC -1- CATALYTIC ACTIVITY: 6-phospho-D-glucono-1,5-lactone + H(2)O = 6-
 CC phospho-D-gluconate.
 CC -1- PATHWAY: Pentose phosphate pathway; second step.
 CC -1- SIMILARITY: BELONGS TO THE GLUCOSAMINE/GALACTOSAMINE-6-PHOSPHATE
 CC ISOMERASE FAMILY. 6-PHOSPHOGLUCONOLACTONASE SUBFAMILY.
 CC -----
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 CC -----
 CC DR EMBL; AF001772; AAD36230.1; -;
 CC DR PIR; F72289; F72289.
 CC DR TIGR; TML154; -;
 CC DR InterPro; IPR006148; Gluc_gal_isom.
 CC DR InterPro; IPR005900; Phosphogluconlac.
 CC Pfam; PF01182; Glucosamine_iso; 1.
 CC DR TIGRfams; TIGR01198; pgl; 1.
 CC KW Hydrolyase; Complete proteome.
 CC SQ SEQUENCE 220 AA; 25325 MW; 9B0FD07E01E60C3 CRC64;
 CC -----
 CC Query Match 48.8%; Score 41; DB 1; Length 220;
 CC Best Local Similarity 40.0%; Pred. No. 14;
 CC Matches 6; Conservative 6; Mismatches 3; Indels 0; Gaps 0;
 Qy 1 QRYGRELRRMSDEFE 15
 Db 113 EKYEREIRSATQDF 127
 RESULT 11
 SNF4_YEAST
 ID SNF4_YEAST STANDARD; PRT; 322 AA.
 AC P12904;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Nuclear protein SNF4 (Regulatory protein CAT3).
 GN SNF4 OR CAT3 OR YGL115W.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 CC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
 OX NCBI_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=89006284; PubMed=3049255;
 RA Schueller H.-J., Entian K.-D.;
 RT "Molecular characterization of yeast regulatory gene CAT3 necessary
 for glucose derepression and nuclear localization of its product.";
 RL Gene 67:247-257(1988).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90097921; PubMed=2481228;
 RA Celenza J.L., Eng F.J., Carlson M.;
 RT "Molecular analysis of the SNF4 gene of Saccharomyces cerevisiae:
 evidence for physical association of the SNF4 protein with the SNF1
 protein kinase.";
 RL Mol. Cell. Biol. 9:5045-5054(1989).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX Lauguin G.;
 RT Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 RL Submitted (JUN-1993) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE OF 30-34 AND 316-322.
 RX MEDLINE=94131988; PubMed=7905477;
 RA Mitchell K.I., Stapleton D., Gao G., House C., Michell B.,
 Katsis F., Witters L.A., Kemp B.E.;
 RT "Mammalian AMP-activated protein kinase shares structural and
 functional homology with the catalytic domain of yeast Snf1 protein
 kinase";
 RL J. Biol. Chem. 269:2361-2364(1994).
 CC -1- FUNCTION: THIS PROTEIN CAUSES EXPRESSION OF GLUCOSE-REPRESSIBLE
 GENES UPON GLUCOSE DEPRIVATION. IT INTERACTS AND HAS FUNCTIONAL
 RELATIONSHIP TO THE PROTEIN-KINASE SNF1.
 CC -1- SUBCELLULAR LOCATION: Nuclear.
 CC -1- SIMILARITY: BELONGS TO THE 5'-AMP-ACTIVATED PROTEIN KINASE, GAMMA
 SUBUNIT FAMILY.
 CC -1- SIMILARITY: Contains 4 CBS domains.
 CC
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 CC
 DR EMBL; M21760; AAA34472.1; -
 DR EMBL; M30470; AAA35061.1; -
 DR EMBL; Z72637; CAA96823.1; -
 DR EMBL; D16506; BAA03958.1; -
 DR PIR; A38906; RBYC3.
 DR SGD; S0003063; SNF4.
 DR GO; GO:0005737; C:cytoplasm; IDA.
 DR GO; GO:0005634; C:nucleus; IDA.
 DR GO; GO:0030295; P:protein kinase activator activity; IGI.
 DR GO; GO:0007031; P:protein kinase organization and biogenesis; IMP.
 DR GO; GO:0006357; P:regulation of transcription from Pol II pro. .; IGI.
 DR InterPro; IPR000644; CBS_domain.
 DR Pfam; PF00571; CBS; 4.
 DR SMART; SM00116; CBS; 4.
 KW Carbohydrate metabolism; Transcription regulation; Nuclear protein;
 Repeat; CBS domain.
 FT DOMAIN 35 89 CBS 1.
 FT DOMAIN 117 175 CBS 2.

FT DOMAIN 192 246 CBS 3.
 FT DOMAIN 259 318 CBS 4.
 SQ SEQUENCE 322 AA; 36401 MW; 51B387E346EB561 CRC64;
 Query Match 48.8%; Score 41; DB 1; Length 322;
 Best Local Similarity 61.5%; Pred. No. 21;
 Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;
 QY 4 GRELRMSDFEG 16
 | | | | | | | | | |
 DB 258 GEALRRSDDFEG 270
 RESULT 12
 SNF4_KLULA
 ID SNF4_KLULA STANDARD; PRT; 328 AA.
 AC Q9P869;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Nuclear protein SNF4.
 GN SNF4.
 OS Kluyveromyces lactis (Yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 CC Saccharomycetales; Saccharomycetaceae; Kluyveromyces.
 OX NCBI_TaxID=28985;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Tomasini L., Ferrero I., Goffrini P.;
 RT "Molecular characterization of KLSNF4 gene";
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: THIS PROTEIN CAUSES EXPRESSION OF GLUCOSE-REPRESSIBLE
 GENES UPON GLUCOSE DEPRIVATION. IT INTERACTS AND HAS FUNCTIONAL
 RELATIONSHIP TO THE PROTEIN-KINASE SNF1 (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Nuclear (By similarity).
 CC -1- SIMILARITY: BELONGS TO THE 5'-AMP-ACTIVATED PROTEIN KINASE, GAMMA
 SUBUNIT FAMILY.
 CC -1- SIMILARITY: Contains 4 CBS domains.
 CC
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 CC
 DR EMBL; AJ277480; CAB89520.1; -
 DR InterPro; IPR000644; CBS_domain.
 DR Pfam; PF00571; CBS; 4.
 DR SMART; SM00116; CBS; 4.
 KW Carbohydrate metabolism; Transcription regulation; Nuclear protein;
 Repeat; CBS domain.
 FT DOMAIN 40 94 CBS 1.
 FT DOMAIN 122 180 CBS 2.
 FT DOMAIN 198 252 CBS 3.
 FT DOMAIN 271 324 CBS 4.
 SQ SEQUENCE 328 AA; 37163 MW; DC9ED3F85E46BD3 CRC64;
 Query Match 48.8%; Score 41; DB 1; Length 328;
 Best Local Similarity 61.5%; Pred. No. 21;
 Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;
 QY 4 GRELRMSDFEG 16
 | | | | | | | | | |
 DB 264 GEALRRSDDFEG 276
 RESULT 13
 PSNR_ARCFU
 ID PSNR_ARCFU STANDARD; PRT; 398 AA.
 AC O28303;
 DT 15-JUL-1998 (Rel. 36, Created)

DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Proteasome-activating nucleotidase (Proteasome regulatory subunit).
 GN PAN OR AF1976.
 OS Archaeoglobus fulgidus.
 OC Archaea; Euryarchaeota; Archaeoglobi; Archaeoglobales;
 OC Archaeoglobaceae; Archaeoglobus.
 OX NCBI_TaxID=2234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=VC-16 / DSM 4304 / ATCC 49558;
 RX MEDLINE=98049343; PubMed=939475;
 RA Klenk H.-P., Clayton R.A., Tomb J.-F., White O., Nelson K.E.,
 RA Ketchum K.A., Dodson R.J., Gwinn M., Hickey E.K., Peterson J.D.,
 RA Richardson D.D., Kariavagale A.R., Graham D.E., Kyrpides N.C.,
 RA Fleischmann R.D., Quackenbush J., Lee N.H., Sutton G.G., Gill S.,
 RA Kirkness E.F., Dougherty B.A., McKenney K., Adams M.D., Loftus B.,
 RA Peterson S., Reich C.I., McNeil L.K., Badger J.H., Glodek A., Zhou L.,
 RA Overbeek R., Gocayne J.D., Weidman J.F., McDonald L., Utterback T.,
 RA Cotton M.D., Spriggs T., Artiach P., Kaine B.P., Sykes S.M.,
 RA Sadow P.W., D'Andrea K.P., Bowman C., Fujii C., Garland S.A.,
 RA Mason T.M., Olsen G.J., Fraser C.M., Smith H.O., Woese C.R.,
 RA Venter J.C.;
 RA "The complete genome sequence of the hyperthermophilic, sulphate-
 RT reducing archaeon Archaeoglobus fulgidus.";
 RT Nature 390:364-370(1997).
 RL Nature 390:364-370(1997).
 CC -!- FUNCTION: Required for the ATP- or CTP-dependent degradation of
 CC proteins, but not small peptides, by the 20S proteasome (By
 CC similarity).
 CC -!- SUBUNIT: Homohexamer (Potential).
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
 CC -!- SIMILARITY: BELONGS TO THE AAA FAMILY OF ATPASES.
 CC -----
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 CC -----
 CC EMBL; AEO00967; AAB89280.1; -;
 DR PIR; G69496; G69496.
 DR TIGR; AF1976; -;
 DR HAMAP; MF_00553; -; 1.
 DR InterPro; IPR005937; 26S_p45.
 DR InterPro; IPR003593; AAA_ATPase.
 DR InterPro; IPR003959; AAA_ATPase_central.
 DR InterPro; IPR003960; AAA_sub.
 DR Pfam; PF00004; AAA; 1.
 DR SMART; SM00382; AAA; 1.
 DR TIGRfam; TIGR01242; 26Sp45; 1.
 DR PROSITE; PS00674; AAA; 1.
 KW Proteasome; ATP-binding; Complete proteome.
 FT NE_BIND 182 189 ATP (POTENTIAL).
 FT SEQUENCE 398 AA; 44964 MW; F3293BB756A646B4 CRC64;
 SQ
 Query Match 48.8%; Score 41; DB 1; Length 398;
 Best Local Similarity 57.1%; Pred. No. 26;
 Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
 QY 2 RYGERLRMSDEFE 15
 || ||:|:| |
 43 RYREVRRLRSEVE 56
 Db
 RESULT 14
 FMRA_CALPA
 ID FMRA_CALPA STANDARD; PRT; 334 AA.
 AC Q01133;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Antho-RFamide neuropeptides precursor.
 OS Calliactis parasitica (Sea anemone).
 OC Eukaryota; Metazoa; Cnidaria; Anthozoa; Zoantharia; Actiniaria;
 OC Nynanthaeae; Hormathiidae; Calliactis.
 OX NCBI_TaxID=6114;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=9117845; PubMed=1706527;
 RA Darner D., Schmutzler C., Diekhoff D., Grimmelikhuijzen C.J.P.;
 RT "Primary structure of the precursor for the sea anemone neuropeptide
 RT Antho-RFamide (<Glu-Gly-Arg-Phe-NEH2>";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:2555-2559(1991).
 CC -!- FUNCTION: NOT KNOWN BUT IT COULD ACT AS A TRANSMITTER AT
 CC NEUROMUSCULAR SYNAPSES.
 CC -!- TISSUE SPECIFICITY: NEURONS ASSOCIATED WITH SMOOTH MUSCLE FIBERS.
 CC -----
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 CC -----
 CC EMBL; M59166; AAA27878.1; -;
 DR PIR; A39172; A39172.
 DR InterPro; IPR002544; FARP.
 DR Pfam; PF01581; FARP; 15.
 KW Neuropeptide; Amidation; Repeat; Signal.
 FT SIGNAL 1 26 POTENTIAL.
 FT PEPTIDE 117 120 ANTHO-RFAMIDE.
 FT PEPTIDE 126 129 ANTHO-RFAMIDE.
 FT PEPTIDE 135 138 ANTHO-RFAMIDE.
 FT PEPTIDE 143 146 ANTHO-RFAMIDE.
 FT PEPTIDE 152 155 ANTHO-RFAMIDE.
 FT PEPTIDE 161 164 ANTHO-RFAMIDE.
 FT PEPTIDE 170 173 ANTHO-RFAMIDE.
 FT PEPTIDE 179 182 ANTHO-RFAMIDE.
 FT PEPTIDE 188 191 ANTHO-RFAMIDE.
 FT PEPTIDE 197 200 ANTHO-RFAMIDE.
 FT PEPTIDE 206 209 ANTHO-RFAMIDE.
 FT PEPTIDE 215 218 ANTHO-RFAMIDE.
 FT PEPTIDE 224 227 ANTHO-RFAMIDE.
 FT PEPTIDE 234 237 ANTHO-RFAMIDE.
 FT PEPTIDE 243 246 ANTHO-RFAMIDE.
 FT PEPTIDE 253 256 ANTHO-RFAMIDE.
 FT PEPTIDE 263 266 ANTHO-RFAMIDE.
 FT PEPTIDE 272 275 ANTHO-RFAMIDE.
 FT PEPTIDE 281 284 ANTHO-RFAMIDE.
 FT MOD_RES 120 120 AMIDATION (G-121 PROVIDE AMIDE GROUP).
 FT MOD_RES 129 129 AMIDATION (G-130 PROVIDE AMIDE GROUP).
 FT MOD_RES 138 138 AMIDATION (G-139 PROVIDE AMIDE GROUP).
 FT MOD_RES 146 146 AMIDATION (G-147 PROVIDE AMIDE GROUP).
 FT MOD_RES 155 155 AMIDATION (G-156 PROVIDE AMIDE GROUP).
 FT MOD_RES 164 164 AMIDATION (G-165 PROVIDE AMIDE GROUP).
 FT MOD_RES 173 173 AMIDATION (G-174 PROVIDE AMIDE GROUP).
 FT MOD_RES 182 182 AMIDATION (G-183 PROVIDE AMIDE GROUP).
 FT MOD_RES 191 191 AMIDATION (G-192 PROVIDE AMIDE GROUP).
 FT MOD_RES 200 200 AMIDATION (G-201 PROVIDE AMIDE GROUP).
 FT MOD_RES 209 209 AMIDATION (G-210 PROVIDE AMIDE GROUP).
 FT MOD_RES 218 218 AMIDATION (G-219 PROVIDE AMIDE GROUP).
 FT MOD_RES 227 227 AMIDATION (G-228 PROVIDE AMIDE GROUP).
 FT MOD_RES 237 237 AMIDATION (G-238 PROVIDE AMIDE GROUP).
 FT MOD_RES 246 246 AMIDATION (G-247 PROVIDE AMIDE GROUP).
 FT MOD_RES 256 256 AMIDATION (G-257 PROVIDE AMIDE GROUP).
 FT MOD_RES 266 266 AMIDATION (G-267 PROVIDE AMIDE GROUP).
 FT MOD_RES 275 275 AMIDATION (G-276 PROVIDE AMIDE GROUP).
 FT MOD_RES 284 284 AMIDATION (G-285 PROVIDE AMIDE GROUP).
 SQ SEQUENCE 334 AA; 39781 MW; 438E182C736EB583 CRC64;
 Query Match 48.2%; Score 40.5; DB 1; Length 334;
 Best Local Similarity 56.2%; Pred. No. 26;
 Matches 9; Conservative 2; Mismatches 4; Indels 1; Gaps 1;

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 ; Search time 17.3714 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-28

Perfect score: 84

Sequence: 1 QRYGRELFRMSDEFEQ 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Lasting first 45 summaries

Database :

SPTREMBL_23:*
1: sp.archaea:*
2: sp.bacteria:*
3: sp.fungi:*
4: sp.human:*
5: sp.invertebrate:*
6: sp.mammal:*
7: sp.mhc:*
8: sp.organelle:*
9: sp.phage:*
10: sp.plant:*
11: sp.podent:*
12: sp.virus:*
13: sp.vertebrate:*
14: sp.unclassified:*
15: sp.virus:*
16: sp.bacteriap:*
17: sp.archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	61	72.6	146	13 Q919N2	Q919n2 brachydanio
2	47	56.0	216	16 Q9KZC5	Q9kzc5 streptomyce
3	45	53.6	230	16 Q8XXS6	Q8xxs6 ralstonia s
4	45	53.6	567	10 Q9FV20	Q9fv20 oryza sativ
5	45	53.6	1696	10 Q8GTN3	Q8gtn3 oryza sativ
6	45	53.6	1748	16 Q989R4	Q989r4 rhizobium l
7	45	53.6	1798	2 Q8KGF8	Q8kgf8 rhizobium l
8	44	52.4	157	16 Q8ES55	Q8es55 oceanobacil
9	44	52.4	564	16 Q9RUK9	Q9ruk9 deinococcus
10	43.5	51.8	1589	2 Q8KUF5	Q8kuf5 actinosynne
11	43	51.2	463	16 Q9HUS7	Q9hus7 pseudomonas
12	43	51.2	674	10 Q93Y86	Q93y86 oryza sativ
13	43	51.2	5635	5 Q9N9N1	Q9n9n1 leishmania
14	42.5	50.6	904	2 Q9KGW3	Q9kgw3 pseudomonas
15	42.5	50.6	909	16 Q8EI34	Q8ei34 shewanella
16	42	50.0	164	17 Q8ZT11	Q8zt11 pyrobaculum

```

17 42 50.0 213 17 Q8TJ31
18 42 50.0 337 3 Q9P5V2
19 42 50.0 375 10 Q94H58
20 42 50.0 425 10 Q8S6J1
21 42 50.0 445 16 Q97I40
22 42 50.0 561 10 Q8LM44
23 42 50.0 581 16 Q8F646
24 42 50.0 606 11 Q8C850
25 42 50.0 606 11 Q8C6N1
26 42 50.0 641 10 Q8GTN7
27 42 50.0 677 10 Q9AYG6
28 42 50.0 804 16 Q8EW78
29 42 50.0 827 10 Q8GTN2
30 42 50.0 829 10 Q9XEM4
31 42 50.0 829 10 Q8GVG4
32 42 50.0 1032 10 Q9LGR7
33 42 50.0 1032 10 Q8LHG3
34 42 50.0 1032 10 Q9LGR6
35 42 50.0 1032 10 Q8SSW6
36 42 50.0 1032 10 Q8RFW0
37 42 50.0 1032 10 Q8GTN5
38 42 50.0 1099 10 Q8W3I5
39 42 50.0 1112 10 Q8LSS6
40 42 50.0 1126 5 Q9VSM2
41 42 50.0 1416 10 Q8S609
42 42 50.0 1429 10 Q94H01
43 42 50.0 1583 10 Q9S7A7
44 42 50.0 1653 5 Q9VSM1
45 42 50.0 1709 5 Q9VSM0

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ALIGNMENTS

RESULT 1

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Q919N2 PRELIMINARY; PRT; 146 AA.
AC Q919N2;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DE 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Bad.
GN BAD.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_taxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20373792; PubMed=10917738;
RA Inohara N., Nunez G.;
RT "Genes with homology to mammalian apoptosis regulators identified in
RT zebrafish.";
RL Cell Death Differ. 7:509-510(2000).
DR EMBL; AF231017; AAF66962.2;
DR HSP; Q92934; I65J.
DR ZFIN; ZDB-GENE-000616-1; bad.
SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;

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Query Match 72.6%; Score 61; DB 13; Length 146;
Best Local Similarity 66.7%; Pred. No. 0.028; 0; Indels 0; Gaps 0;
Matches 10; Conservative 5; Mismatches 5;

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QY 1 QRYGRELFRMSDEFE 15
Db 93 KKYGQQLRMSDEFD 107
:::|||||:

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RESULT 2

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Q9KZC5 PRELIMINARY; PRT; 216 AA.
AC Q9KZC5;

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DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Hypothetical protein SC06964.
GN SC06964 OR SC6F7.17C.
OS Streptomyces coelicolor.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=1902;
RN [1]
SEQUENCE FROM N.A.
RP STRAIN=A3(2);
RC Saunders D.C., Harris D.;
RA Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
SEQUENCE FROM N.A.
RP STRAIN=A3(2);
RC Thomson N.R., Parkhill J., Barrell B.G., Rajandream M.A.;
RA Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
SEQUENCE FROM N.A.
RP STRAIN=A3(2);
RC MEDLINE=97000351; PubMed=8843436;
RX Redenbach M., Kieser H.M., Denapalte D., Eichner A., Cuillum J.,
RA Kinashi H., Hopwood D.A.;
RT "A set of ordered cosmids and a detailed genetic and physical map for
RL the 8 Mb Streptomyces coelicolor A3(2) chromosome.";
RN Mol. Microbiol. 21:77-96(1996).
RN [4]
SEQUENCE FROM N.A.
RP STRAIN=A3(2) / M145;
RC MEDLINE=21996410; PubMed=12000953;
RX Bentley S.D., Chater K.F., Cerdeno-Tarraga A.-M., Challis G.L.,
RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,
RA Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,
RA Cronin A., Fraser A., Goble A., Hidalgo J., Hornsby T., Howarth S.,
RA Huang C.-H., Kaser T., Larke L., Murphy L., Oliver K., O'Neil S.,
RA Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,
RA Seeger K., Saunders D., Sharp S., Squares S., Taylor K.,
RA Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,
RA Hopwood D.A.;
RT "Complete genome sequence of the model actinomycete Streptomyces
RL coelicolor A3(2).";
RN Nature 417:141-147(2002).
DR EMBL: AL939129; CAB89025.1; -.
DR InterPro: IPR003265; Endo_3c.
DR Pfam: PF00730; Hnr-GPD; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 216 AA; 23810 MW; DDD7C717D60F7AA7 CRC64;

Query Match 56.0%; Score 47; DB 16; Length 216;
Best Local Similarity 50.0%; Pred. No. 8.7;
Matches 8; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDFEG 16
Db 109 ERWGGDLRLRDEADG 124

RESULT 3
Q8XXS6 PRELIMINARY; PRT; 230 AA.
AC Q8XXS6;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Probable ATP-binding ABC transporter protein.
GN RSC2037 OR RS03602.
OS Ralstonia solanacearum (Pseudomonas solanacearum).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Ralstoniaceae; Ralstonia.
OX NCBI_TaxID=305;
RN [1]

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RP SEQUENCE FROM N.A.
RC STRAIN=GMI1000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
RA Arlat M., Billault A., Brottier P., Camus J.C., Cattolico L.,
RA Chandler M., Choisme N., Claudel-Renard C., Cunnac S., Demange N.,
RA Gaspin C., Lavie M., Moisan A., Robert C., Saurin W., Schiex T.,
RA Sigauier P., Thebault P., Whalen M., Wincker P., Levy M.,
RA Weissenbach J., Boucher C.A.;
RT "Genome sequence of the plant pathogen Ralstonia solanacearum.";
RL Nature 415:497-502(2002).
DR EMBL: AL646067; CAD15739.1; -.
DR InterPro: IPR003593; AAA_Atpase.
DR InterPro: IPR003439; ABC_transporter.
DR Pfam: PF00005; ABC_tran; 1.
DR ProDom: PD000006; ABC_transporter; 1.
DR SMART: SM00382; AAA; 1.
KW Complete proteome.
SQ SEQUENCE 230 AA; 25231 MW; 7C3FDA1E7A19A2A4 CRC64;

Query Match 53.6%; Score 45; DB 16; Length 230;
Best Local Similarity 64.3%; Pred. No. 20;
Matches 9; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDFEG 14
Db 168 QEIGTLLRLVDF 181

RESULT 4
Q9FV20 PRELIMINARY; PRT; 567 AA.
ID Q9FV20;
AC Q9FV20;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein.
OS Oryza sativa (indica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=39946;
RN [1]
SEQUENCE FROM N.A.
RP STRAIN=cv. Zhaiyueqing 8; TRANSPOSON-unspecific; TISSUE=Shoot;
RA Yu F., Zhang A., Zhang F., Chen S.;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF259776; AAG15480.1; -.
DR Gramene; Q9FV20; -.
DR InterPro: IPR002156; RNaseH.
DR InterPro: IPR000477; RVase.
DR Pfam: PF00075; rnaseh; 1.
DR Pfam: PF00078; rvt; 1.
KW RNA-directed DNA polymerase; Transferase.
SQ SEQUENCE 567 AA; 64419 MW; BD4F47AB33A685BE CRC64;

Query Match 53.6%; Score 45; DB 10; Length 567;
Best Local Similarity 50.0%; Pred. No. 52;
Matches 7; Conservative 5; Mismatches 2; Indels 0; Gaps 0;

QY 3 YGRELRRMSDFEG 16
Db 547 YRQEVKLEDKFEG 560

RESULT 5
Q8GTN3 PRELIMINARY; PRT; 1696 AA.
ID Q8GTN3;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Putative GAG-POL precursor.

```

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GN OSJNBA0041114.29.
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliopsida; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Liu J., Gansberger K., Jones K.M.,
RA Overton II L.B., Tsitrin T., Kim M.M., Bera J.J., Jin S.S.,
RA Fadrosch D.W., Tallon L.J., Koo H., Zismann V., Hsiao J., Blunt S.,
RA Vanaken S.S., Riedmuller S.B., Utterback T.T., Feldhlyum T.V.,
RA Yang Q.Q., Haas B.J., Suh B.B., Peterson J.J., Quackenbush J.,
RA White O., Salzberg S.L., Fraser C.M.;
RT "Oryza sativa chromosome 10 BAC OSJNBA0041114 genomic sequence.";
RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC099042; AA08247.1;
SQ SEQUENCE 1696 AA; 190501 MW; 516B35CFDFA2F068 CRC64;

Query Match 53.6%; Score 45; DB 10; Length 1696;
Best Local Similarity 50.0%; Pred. No. 1.7e+02;
Matches 7; Conservative 5; Mismatches 2; Indels 0; Gaps 0;

QY 3 YGRELRRMSDEFE 16
DB 1341 YQEVKLEDKFEG 1354

RESULT 6
Q989R4 PRELIMINARY; PRT; 1748 AA.
AC Q989R4;
DT 01-OCT-2001 (TRENBLrel. 18, Created)
DT 01-OCT-2001 (TRENBLrel. 18, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Hypothetical protein mlr6316.
GN MLR6316.
OS Rhizobium loti (Mesorhizobium loti).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Phyllobacteriaceae; Mesorhizobium.
OX NCBI_TaxID=381;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MAFF303099;
RA Kaneko T., Nakamura Y., Sato S., Asamizu E., Kato T., Sasamoto S.,
RA Watanabe A., Idesawa K., Ishikawa A., Kawashima K., Kimura T.,
RA Kishida Y., Kiyokawa C., Kohara M., Matsumoto M., Matsuno A.,
RA Mochizuki Y., Nakayama S., Nakazaki N., Shimpo S., Sugimoto M.,
RA Takeuchi C., Yamada M., Tabata S.;
RT "Complete genome structure of the nitrogen-fixing symbiotic bacterium
RT Mesorhizobium loti";
RL DNA Res. 7:331-338(2000).
DR EMBL; AF003008; BAB52630.1;
DR InterPro; IPR003653; SUMO.protease.
DR Pfam; PF02902; Peptidase_C48; 1.
DR PROSITE; PS50600; ULP_PROTEASE; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 1748 AA; 190454 MW; 2A9CE7B92C354341 CRC64;

Query Match 53.6%; Score 45; DB 16; Length 1748;
Best Local Similarity 69.2%; Pred. No. 1.7e+02;
Matches 9; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRELRRMSDEFE 15
DB 562 YGELRRFSEELE 574

RESULT 7
Q8KGy8 PRELIMINARY; PRT; 1798 AA.
ID Q8KGy8
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Q8KGy8;
AC 01-OCT-2002 (TRENBLrel. 22, Created)
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Hypothetical protein.
GN MSI059.
OS Rhizobium loti (Mesorhizobium loti).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Phyllobacteriaceae; Mesorhizobium.
OX NCBI_TaxID=381;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=R7A;
RA MEDLINE=21599272; PubMed=12003951;
RA Sullivan J.T., Tzibiatowski J.R., Cruickshank R.W., Gouzy J.,
RA Brown S.D., Elliot R.M., Fleetwood D.J., McCallum N.G., Rossbach U.,
RA Stuart G.S., Weaver J.E., Webby R.J., de Bruijn F.J., Ronson C.W.;
RT "Comparative sequence analysis of the symbiosis island of
RT Mesorhizobium loti strain R7A.";
RL J. Bacteriol. 184:3086-3095(2002).
DR EMBL; AL672112; CAD31464.1;
DR InterPro; IPR003653; SUMO.protease.
DR Pfam; PF02902; Peptidase_C48; 1.
DR PROSITE; PS50600; ULP_PROTEASE; 1.
KW Hypothetical protein.
SQ SEQUENCE 1798 AA; 196214 MW; A750A49B8C0D581B CRC64;

Query Match 53.6%; Score 45; DB 2; Length 1798;
Best Local Similarity 69.2%; Pred. No. 1.8e+02;
Matches 9; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRELRRMSDEFE 15
DB 569 YGELRRFSEELE 581

RESULT 8
Q8ES55 PRELIMINARY; PRT; 157 AA.
AC Q8ES55;
DT 01-MAR-2003 (TRENBLrel. 23, Created)
DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Hypothetical conserved protein.
GN OB0788.
OS Oceanobacillus ihyenssis.
OC Bacteria; Firmicutes; Bacillales; Oceanobacillus.
OX NCBI_TaxID=182710;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=HTE831 / DSM 14371 / JCM 11309;
RX MEDLINE=22220767; PubMed=12235376;
RA Takami H., Takaki Y., Uchiyama I.;
RT "Genome sequence of Oceanobacillus ihyenssis isolated from the Iheya
RT Ridge and its unexpected adaptive capabilities to extreme
RT environments.";
RL Nucleic Acids Res. 30:3927-3935(2002).
DR EMBL; AF004595; BAC12744.1;
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 157 AA; 17809 MW; 128229251596C18C CRC64;

Query Match 52.4%; Score 44; DB 16; Length 157;
Best Local Similarity 50.0%; Pred. No. 19;
Matches 7; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRELRRMSDEFE 16
DB 126 FSELKRIGEEFSG 139

RESULT 9
Q9RUK9 PRELIMINARY; PRT; 564 AA.
ID Q9RUK9
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AC Q9RUK9;
DT 01-MAY-2000 (TRENBLrel. 13, Created)
DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Glycosyl hydrolase, family 13.
GN DR1375.
OS Deinococcus radiodurans.
OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
OC Deinococcaceae; Deinococcus.
OX NCBI_TaxID=1299;
FN [1]
RP SEQUENCE FROM N.A.
PC STRAIN=R1;
EX MEDLINE=20036896; PubMed=10567266;
RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
RA Vamathevan J.J., Lam P., McDonald L., Utterback T., Zalewski C.,
RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RT "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1.";
RL Science 286:1571-1577(1999).
DR EMBL; AE001983; AAF10944.1; -.
DR HSP; P21332; 1UOK.
DR TIGR; DR1375; -.
DR InterPro; IPR006047; Alpha_amyl_cat.
DR InterPro; IPR006589; Alp_amyl_cat_sub.
DR Pfam; PF001128; alpha-amylase; 1.
DR SMART; SM00642; Aamy; 1.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 564 AA; 63667 MW; B8F50B9B0DFC8D51 CRC64;

Query Match 52.4%; Score 44; DB 16; Length 564;
Best Local Similarity 61.5%; Pred. No. 76;
Matches 8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 YGELRRMSDEFE 15
DB 283 YVEMRRVIDEFD 295

RESULT 10
O8KUF5
ID O8KUF5 PRELIMINARY; PRT; 1589 AA.
AC O8KUF5;
DT 01-OCT-2002 (TRENBLrel. 22, Created)
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Polyketide synthase.
GN ASWG..
OS Actinosynnema pretiosum (subsp. aurantium).
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Pseudonocardineae; Actinosynnemataceae; Actinosynnema.
OX NCBI_TaxID=42198;
FN [1]
RP SEQUENCE FROM N.A.
PC STRAIN=ATCC 31565;
EX MEDLINE=22056096; PubMed=12060743;
RA Yu T.W., Bai L., Clade D., Hoffmann D., Toelzer S., Trinh K.O., Xu J.,
RA Moss S.J., Leistner E., Floss H.G.;
RT "The biosynthetic gene cluster of the maytansinoid antitumor agent
RT ansamitocin from Actinosynnema pretiosum.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:7968-7973(2002).
DR EMBL; AF453501; AAM54077.1; -.
DR InterPro; IPR001227; Ac transferase.
DR InterPro; IPR000794; Ketoacyl-synt.
DR InterPro; IPR006162; Ppantne_attach.
DR InterPro; IPR006163; Pp_bind.
DR Pfam; PF00698; Acyl_transf; 1.
DR Pfam; PF00109; ketoacyl-synt; 1.
DR Pfam; PF02801; ketoacyl-synt_C; 1.

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DR Pfam; PF00550; pp-binding; 1.
DR PROSITE; PS00075; ACP_DOMAIN; 1.
DR PROSITE; PS00606; B_KETOACYL_SYNTHASE; 1.
DR PROSITE; PS00012; PHOSPHOPANTHETHEINE; 1.
KW Phosphopantetheine.
SQ SEQUENCE 1589 AA; 164712 MW; 5067CF9A772A525F CRC64;

Query Match 51.8%; Score 43.5; DB 2; Length 1589;
Best Local Similarity 56.2%; Pred. No. 2.7e+02;
Matches 9; Conservative 3; Mismatches 3; Indels 1; Gaps 1;

QY 1 QYGLRRMSDEFE 16
DB 161 QNYGNR-RVAAEFEG 175

RESULT 11
Q9HUS7
ID Q9HUS7 PRELIMINARY; PRT; 463 AA.
AC Q9HUS7;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Probable two-component sensor.
GN PA4886.
OS Pseudomonas aeruginosa.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=287;
FN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 15692 / PA01;
EX MEDLINE=20437337; PubMed=10984043;
RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
RA Hickey M.J., Brinkman F.S.L., Huffnagle W.O., Kowalik D.J., Lagrou M.,
RA Garber R.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
RA Brody L.L., Gaulty S., Tolentino E., Westbrook-Wadman S., Yuan Y.,
RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.;
RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an
RT opportunistic pathogen.";
RL Nature 406:959-964(2000).
CC -!- SIMILARITY: TO PROKARYOTE SENSORY TRANSDUCTION PROTEINS.
DR EMBL; AE004901; AAG08271.1; -.
DR HSP; P02933; 1BYD.
DR InterPro; IPR003594; ATPbind_ATPase.
DR InterPro; IPR004358; Bact_sens_pr_C.
DR InterPro; IPR003660; HAMP.
DR InterPro; IPR003661; His_KinA.
DR InterPro; IPR005467; His_Kinase.
DR InterPro; IPR000005; HTHArac.
DR InterPro; IPR006290; Metal_his_kin.
DR Pfam; PF00672; HAMP; 1.
DR Pfam; PF02518; HATPase_c; 1.
DR Pfam; PF00512; Hiska; 1.
DR PRINTS; PR00344; BCTRLSENSOR.
DR SMART; SM00304; HAMP; 1.
DR SMART; SM00387; HATPase_c; 1.
DR SMART; SM00388; Hiska; 1.
DR TIGRFAMS; TIGR01386; czts_sils_cops; 1.
DR PROSITE; PS0109; HIS_KIN; 1.
KW Kinase; Phosphorylation; Sensory transduction; Transferase;
KW Complete proteome.
SQ SEQUENCE 463 AA; 50946 MW; 97FF19C2CF38006C CRC64;

Query Match 51.2%; Score 43; DB 16; Length 463;
Best Local Similarity 61.5%; Pred. No. 90;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 4 GRELRRMSDEFE 16
DB 321 GEELRRQAEFEFEG 333

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DR SMART: SM00382; AAA: 4.
DR PROSITE; PS00639; THIOL_PROTEASE_HIS; 1.
KW ATP-binding.
SQ SEQUENCE 5635 AA; 620050 MW; 6A9AE81A9B14641 CRC64;

Query Match          51.2%; Score 43; DB 5; Length 5635;
Best Local Similarity 72.7%; Pred. No. 1.3e+03;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 QRYGRELRRMS 11
Db 1535 QRTGRDLRW 1545

RESULT 14
Q9KGW3 PRELIMINARY; PRT; 904 AA.
ID Q9KGW3 AC Q9KGW3
RC STRAIN=cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Moffat K.S., Hill J.N., Gansberger K.,
RA Brenner M., Burgess S., Hance M., Shvartsbeyn M., Tsitrin T.,
RA Riggs F., Hsiao J., Zismann V., Blunt S., Pal G., VanAken S.E.,
RA Uterback T.R., Feldblum T.V., Quackenbush J., Salzberg S.L.,
RA White O., Fraser C.M.;
RT "Oryza sativa chromosome 3 BAC OSJNBa0026A15 genomic sequence.";
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC084404; AAK50597.1; -.
DR Gramene; Q93Y86; -.
DR InterPro; IPR002156; RNaseH.
DR InterPro; IPR001584; Rve.
DR Pfam; PF00075; rnaseH; 1.
DR Pfam; PF00665; rve; 1.
SQ SEQUENCE 674 AA; 75473 MW; 9FA929B1426E725E CRC64;

Query Match          51.2%; Score 43; DB 10; Length 674;
Best Local Similarity 50.0%; Pred. No. 1.3e+02;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFG 16
Db 258 EAYVREYRRMRNFDG 273

RESULT 13
Q9NSN1 PRELIMINARY; PRT; 5635 AA.
ID Q9NSN1 AC Q9NSN1
RC STRAIN=Friedlin;
RA Hilbert H., Wedler H., Wedler E., Dueterhoeft A., Ivens A.C.,
RA Quail M., Rajandream M.A., Barrell B.G.;
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
SQ SEQUENCE FROM N.A.
RC STRAIN=Friedlin;
RX MEDLINE=98146435; PubMed=9477341;
RA Ivens A.C., Lewis S.M., Bagherzadeh A., Zhang L., Chan H.M.,
RA Smith D.F.;
RT "A physical map of the Leishmania major Friedlin genome.";
RL Genome Res. 8:135-145(1998).
DR EMBL; AL359781; CAB95305.1; -.
DR InterPro; IPR003593; AAA_AtpPase.
DR InterPro; IPR004273; Dynein_heavy.
DR InterPro; IPR000169; SHprot_aseite.
DR Pfam; PF03028; Dynein_heavy; 1.

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DR SMART: SM00382; AAA: 4.
DR PROSITE; PS00639; THIOL_PROTEASE_HIS; 1.
KW ATP-binding.
SQ SEQUENCE 5635 AA; 620050 MW; 6A9AE81A9B14641 CRC64;

Query Match          51.2%; Score 43; DB 5; Length 5635;
Best Local Similarity 72.7%; Pred. No. 1.3e+03;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 QRYGRELRRMS 11
Db 1535 QRTGRDLRW 1545

RESULT 14
Q9KGW3 PRELIMINARY; PRT; 904 AA.
ID Q9KGW3 AC Q9KGW3
RC STRAIN=WCS365;
RA Camacho Carvajal M.M., Lugtenberg B.J.J., Bloemberg G.V.;
RT "Characterization of NADH dehydrogenases of Pseudomonas fluorescens
RT WCS365 and their role in competitive root colonisation.";
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF281148; AAF97803.1; -.
DR InterPro; IPR000283; Complex1_75K.
DR InterPro; IPR001041; Ferredoxin.
DR InterPro; IPR006656; Molybdopterin.
DR InterPro; IPR006963; Molybdop_Fe4S4.
DR Pfam; PF00111; fer2; 1.
DR Pfam; PF00384; molybdopterin; 1.
DR Pfam; PF04879; Molybdop_Fe4S4; 1.
DR PROSITE; PS00641; COMPLEX1_75K_1; 1.
DR PROSITE; PS00642; COMPLEX1_75K_2; 1.
DR PROSITE; PS00643; COMPLEX1_75K_3; 1.
KW Iron; Iron-sulfur.
SQ SEQUENCE 904 AA; 98157 MW; C25E86C6D4DFA457 CRC64;

Query Match          50.6%; Score 42.5; DB 2; Length 904;
Best Local Similarity 56.2%; Pred. No. 2.2e+02;
Matches 9; Conservative 3; Mismatches 3; Indels 1; Gaps 1;

QY 1 QRYGRELRRMSDEFG 16
Db 241 ERYG-ELRRNRFNG 255

RESULT 15
Q8EI34 PRELIMINARY; PRT; 909 AA.
ID Q8EI34 AC Q8EI34
RC STRAIN=cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Moffat K.S., Hill J.N., Gansberger K.,
RA Brenner M., Burgess S., Hance M., Shvartsbeyn M., Tsitrin T.,
RA Riggs F., Hsiao J., Zismann V., Blunt S., Pal G., VanAken S.E.,
RA Uterback T.R., Feldblum T.V., Quackenbush J., Salzberg S.L.,
RA White O., Fraser C.M.;
RT "Oryza sativa chromosome 3 BAC OSJNBa0026A15 genomic sequence.";
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC084404; AAK50597.1; -.
DR Gramene; Q93Y86; -.
DR InterPro; IPR002156; RNaseH.
DR InterPro; IPR001584; Rve.
DR Pfam; PF00075; rnaseH; 1.
DR Pfam; PF00665; rve; 1.
SQ SEQUENCE 674 AA; 75473 MW; 9FA929B1426E725E CRC64;

Query Match          51.2%; Score 43; DB 10; Length 674;
Best Local Similarity 50.0%; Pred. No. 1.3e+02;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFG 16
Db 258 EAYVREYRRMRNFDG 273

RESULT 13
Q9NSN1 PRELIMINARY; PRT; 5635 AA.
ID Q9NSN1 AC Q9NSN1
RC STRAIN=Friedlin;
RA Hilbert H., Wedler H., Wedler E., Dueterhoeft A., Ivens A.C.,
RA Quail M., Rajandream M.A., Barrell B.G.;
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
SQ SEQUENCE FROM N.A.
RC STRAIN=Friedlin;
RX MEDLINE=98146435; PubMed=9477341;
RA Ivens A.C., Lewis S.M., Bagherzadeh A., Zhang L., Chan H.M.,
RA Smith D.F.;
RT "A physical map of the Leishmania major Friedlin genome.";
RL Genome Res. 8:135-145(1998).
DR EMBL; AL359781; CAB95305.1; -.
DR InterPro; IPR003593; AAA_AtpPase.
DR InterPro; IPR004273; Dynein_heavy.
DR InterPro; IPR000169; SHprot_aseite.
DR Pfam; PF03028; Dynein_heavy; 1.

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RX MEDLINE=22297686; PubMed=12368813;
 RA Heidelberg J.F., Paulsen I.T., Nelson K.E., Gaidos E.J., Nelson W.C.,
 RA Read T.D., Eisen J.A., Seshadri R., Ward N., Methe B., Clayton R.A.,
 RA Meyer T., Tsapin A., Scott J., Beanan M., Brinkac L., Daugherty S.,
 RA DeBoy R.T., Dodson R.J., Durkin A.S., Haft D.H., Kolonay J.F.,
 RA Madupu R., Peterson J.D., Umayam L.A., White O., Wolf A.M.,
 RA Vamathevan J., Weidman J., Impraim M., Lee K., Berry K., Lee C.,
 RA Mueller J., Khouri H., Gill J., Utterback T.R., McDonald L.A.,
 RA Feldblyum T.V., Smith H.O., Venter J.C., Nelson K.H., Fraser C.M.;
 RT "Genome sequence of the dissimilatory metal ion-reducing bacterium
 RT *Shewanella oneidensis*.";
 RL Nat. Biotechnol. 20:1118-1123(2002).
 DR EMBL; AE015546; AAN54089.1; -;
 DR TIGR; SO1016; -;
 KW Complete proteome.
 SQ SEQUENCE 909 AA; 100079 MW; F71859F385BDCAC8 CRC64;

Query Match 50.6%; Score 42.5; DB 16; Length 909;
 Best Local Similarity 56.2%; Pred. No. 2.2e+02;
 Matches 9; Conservative 3; Mismatches 3; Indels 1; Gaps 1;

QY 1 QRYGRLRMSDEPEG 16
 :||| ||||: ||
 Db 241 ERYG-ELRRERFRG 255

Search completed: September 15, 2003, 17:25:48
 Job time : 19.3714 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:01 ; Search time 22.6286 Seconds
(without alignments)
112.231 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRELRRMSDEFVD 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 24: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	83	100.0	16	20	AA1980
2	83	100.0	16	20	AA1981
3	83	100.0	20	23	AA1982
4	83	100.0	20	23	AA1983
5	83	100.0	21	23	AA1984
6	83	100.0	21	23	AA1985
7	83	100.0	22	23	AA1986
8	83	100.0	23	23	AA1987
9	83	100.0	24	23	AA1988

10	83	100.0	24	23	AA1989
11	83	100.0	24	23	AA1990
12	83	100.0	25	23	AA1991
13	83	100.0	25	23	AA1992
14	83	100.0	25	23	AA1993
15	83	100.0	25	23	AA1994
16	83	100.0	25	23	AA1995
17	83	100.0	25	23	AA1996
18	83	100.0	25	23	AA1997
19	83	100.0	25	23	AA1998
20	83	100.0	25	23	AA1999
21	83	100.0	25	23	AA2000
22	83	100.0	25	23	AA2001
23	83	100.0	25	23	AA2002
24	83	100.0	25	23	AA2003
25	83	100.0	25	23	AA2004
26	83	100.0	25	23	AA2005
27	83	100.0	25	23	AA2006
28	83	100.0	25	23	AA2007
29	83	100.0	25	23	AA2008
30	83	100.0	25	23	AA2009
31	83	100.0	25	23	AA2010
32	83	100.0	25	23	AA2011
33	83	100.0	25	23	AA2012
34	83	100.0	25	23	AA2013
35	83	100.0	25	23	AA2014
36	83	100.0	25	23	AA2015
37	83	100.0	25	23	AA2016
38	83	100.0	25	23	AA2017
39	83	100.0	25	23	AA2018
40	83	100.0	26	21	AA2019
41	83	100.0	26	22	AA2020
42	83	100.0	166	18	AA2021
43	83	100.0	168	19	AA2022
44	83	100.0	168	21	AA2023
45	83	100.0	168	22	AA2024

ALIGNMENTS

RESULT 1
AA1980
ID AA1980 standard; peptide; 16 AA.
AC AA1980;
DT 02-JUL-1999 (first entry)
DE Human BAD BH3 domain.
KW BH3 domain; cell death agonist; bcl homology domain; Bcl-2 family;
KW apoptosis promoter; cancer cell; virus infected cell; inflammation;
KW autoantibody producing cell; cancer; lymphoproliferative condition;
KW arthritis; autoimmune disease; therapy.
XX Homo sapiens.
XX WO9916787-A1.
XX 08-APR-1999.
XX 22-SEP-1998; 98WO-US19765.
XX 07-OCT-1997; 97US-0946039.
XX 26-SEP-1997; 97US-0060133.
XX (UNTW) UNIV WASHINGTON.
XX Korsmeyer SJ;
XX WPI; 1999-2550508/21.

PT Bcl homology domain 3 polypeptide
XX Example 1; Fig 4; 104pp; English.

XX This sequence represents the BH3 domain of human BAd.
CC The invention relates to a bcl homology domain 3 (BH3 domain),
CC derived from a proapoptotic member of the BCL-2 family. The
CC BH3 polypeptide can be used in a method for promoting apoptosis in a
CC target cell, especially where the cell is a cancer cell a virus infected
CC cell or an autobody producing cell. The BH3 polypeptide can be used
CC in therapeutic compositions for treating disease including cancer, other
CC lymphoproliferative conditions, arthritis, inflammation, and autoimmune
CC diseases, which may result from the down regulation of cell death
CC regulation.

XX Sequence 16 AA;

Query Match 100.0%; Score 83; DB 20; Length 16;
Best Local Similarity 100.0%; Pred. No. 1.3e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDFVD 16
DB 1 QRYGRLRMSDFVD 16

RESULT 2
AAB37029
ID AAB37029 standard; peptide; 16 AA.

XX AAB37029;
AC
DT 28-FEB-2001 (first entry)

XX Bcl2 polypeptide BH3 domain peptide #29.

XX Cytostatic; neuroprotective; anti-HIV; virucide; cerebroprotective;
KW cardiant; Bcl-2 superfamily; BH3 domain; cell death agonist; BAd;
KW apoptosis modulation; B cell lymphoma/leukemia 2; cancer; prostate;
KW colorectal; gastric; non-small lung; renal; thyroid; neuroblastoma;
KW melanoma; lymphocytic leukemia; neurodegenerative disorder; AIDS;
KW stroke; myocardial infarction.

XX Homo sapiens.

XX WO2000059526-A1.

XX 12-OCT-2000.

XX 06-APR-2000; 2000WO-US09352.

XX 07-APR-1999; 99US-0128202.

XX (UYJE-) UNIV JEFFERSON THOMAS.

XX Huang Z, Wang J, Zhang Z, Shan S, Lu Z;

XX WPI; 2000-679325/66.

XX New peptide conjugates for modulating apoptosis or for inhibiting B
PT cell lymphoma/leukemia 2 (Bcl-2) function, especially useful for
PT treating neurodegenerative disorders, stroke, or cancer

XX Claim 18; Page 18; 74pp; English.

XX The invention relates to a peptide conjugate having the formula:
CC (R-X)n-peptide where n = 1-10; x = C=O, when the R-x group is attached
CC to the N-terminus of the peptide, or a side chain of the peptide where
CC the functional group of the side chain is NH2 or OH; or x = O or NH.
CC when the R-X group is attached to the C-terminus of the peptide, or a
CC side chain of the peptide, where the side chain functional group is COOH
CC or CONH2; and R = 2-18C alkyl or alkoxy, 2-14C alkylphenyl containing one
CC or two double bonds, cyclohexyl, cyclopentyl, cyclohexyl optionally

CC monosubstituted with a 1-5C straight or branched chain alkyl group,
CC phenyl optionally monosubstituted with a 1-5C straight or branched chain
CC alkyl group, or benzyl. The peptides AAB37001-B37058 represent examples
CC of the peptide portion of the conjugate. The peptides represent analogues
CC of a Bcl-2 superfamily polypeptide corresponding to amino acids 72-97 of
CC the BH3 domain of the cell death agonist BAd. The peptide conjugate is
CC useful for modulating apoptosis in the cells of a subject, or for
CC reversing B cell lymphoma/leukemia 2 (Bcl-2)-mediated blockage of
CC apoptosis in cancer cells. It is also useful for inhibiting Bcl-2
CC function. In particular, the peptide conjugate is useful for treating a
CC subject afflicted with a cancer characterized by cancer cells that
CC express Bcl-2. The cancer includes prostate, colorectal, gastric,
CC non-small lung, renal or thyroid cancers, neuroblastoma, melanoma, or
CC acute or chronic lymphocytic and non-lymphocytic leukemia. The peptide
CC conjugate is also useful for treating disorders characterized by
CC increased apoptosis, e.g. neurodegenerative disorders, acquired
CC immunodeficiency syndrome (AIDS), stroke or myocardial infarction.

XX Sequence 16 AA;

Query Match 100.0%; Score 83; DB 21; Length 16;
Best Local Similarity 100.0%; Pred. No. 1.3e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDFVD 16
DB 1 QRYGRLRMSDFVD 16

RESULT 3
ABG78499

XX ABG78499 standard; Peptide; 20 AA.

XX AC ABG78499;

XX 15-NOV-2002 (first entry)

XX Mutant Bcl2 competitive binding assay peptide #16.

XX Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; mutain.

XX Homo sapiens.

XX Synthetic.

XX WO2002040530-A2.

XX 23-MAY-2002.

XX 15-NOV-2001; 2001WO-US45693.

XX 20-NOV-2000; 2000US-0716395.

XX (ABBO) ABBOTT LAB.

XX Fesik SW, Petros AM, Yoon H, Nettesheim DG;

XX WPI; 2002-490141/52.

XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
PT useful in biological assays to identify substances that block the
PT ability of Bcl-2 to inhibit programmed cell death or apoptosis

XX Example 2; Page 17; 36pp; English.

XX This invention relates to a novel mutant protein which is derived from
CC a wild type human Bcl-2 protein. The mutant is created by replacing a
CC sequence of amino acid residues comprising a flexible loop from the wild
CC type Bcl-2 protein with an amino acid sequence comprising at least two
CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
CC shown in the specification. The invention also comprises an assay for
CC identifying substances that bind to the Bcl-2 protein. The protein
CC sequences of the invention are useful in biological assays to identify
CC substances that block the ability of Bcl-2 to inhibit programmed cell

CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.

SQ Sequence 20 AA;

Query Match 100.0%; Score 83; DB 23; Length 20;
 Best Local Similarity 100.0%; Pred. No. 1.7e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRLRMSDFVD 16
 |||||
 Db 1 QRYGRLRMSDFVD 16

RESULT 4

AAU78626
 ID AAU78626 standard; Peptide: 20 AA.

XX AC AAU78626;

DT 18-JUN-2002 (first entry)

XX Human Bad peptide #26 which binds to a member of the Bcl-2 family.

XX Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
 KW ischemic injury; suppressor; BH3 domain.

XX Homo sapiens.

OS WC200220568-A2.

XX PD 14-MAR-2002.

XX PF 04-SEP-2001; 2001WO-US27410.

XX PR 06-SEP-2000; 2000US-0656399.

XX (ABBO) ABBOTT LAB.

XX Pi Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
 Pi Nettesheim DG, Swift KM, Matayoshi E, Zhang H;

DR WPI; 2002-292254/33.

XX New derivatives of Bad peptide, useful for identifying compounds that
 PT bind to Bcl-2 proteins, potential agents for treating cancer and
 PT degenerative disease -

XX Example 1; Page 14; 31pp; English.

XX The present invention relates to new peptides that are derived from a
 CC wild-type human Bad peptide and are able to bind to a member of the
 CC Bcl-2 protein family. The peptides are useful, when labelled, in
 CC competitive/displacement assays for identifying substances that bind to
 CC members of the Bcl-2 family and may induce or suppress apoptosis so are
 CC potentially useful for treating cancer (inducers) or degenerative
 CC diseases or ischemic injury (suppressors). The peptides of the invention
 CC have high helix propensity, maintain the contacts of the wild-type Bad
 CC peptide and, compared with the Bad peptide, may have better physical
 CC properties, particularly solubility. The present sequence represents one
 CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
 CC from the BH3 domain of the human wild-type Bad peptide.

XX Sequence 20 AA;

Query Match 100.0%; Score 83; DB 23; Length 20;
 Best Local Similarity 100.0%; Pred. No. 1.7e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRLRMSDFVD 16
 |||||
 Db 1 QRYGRLRMSDFVD 16

RESULT 5

ABG78500
 ID ABG78500 standard; Peptide: 21 AA.

XX AC ABG78500;

DT 15-NOV-2002 (first entry)

XX Mutant Bcl2 competitive binding assay peptide #17.

XX Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; mutin.

OS Homo sapiens.

OS Synthetic.

XX PN WO200240530-A2.

XX PD 23-MAY-2002.

XX PF 15-NOV-2001; 2001WO-US45693.

XX PR 20-NOV-2000; 2000US-0716395.

XX PA (ABBO) ABBOTT LAB.

XX Pi Fesik SW, Petros AM, Yoon H, Nettesheim DG;

XX WPI; 2002-490141/52.

XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
 PT useful in biological assays to identify substances that block the
 PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -

XX Example 2; Page 17; 36pp; English.

XX This invention relates to a novel mutant protein which is derived from
 CC a wild type human Bcl-2 protein. The mutant is created by replacing a
 CC sequence of amino acid residues comprising a flexible loop from the wild
 CC type Bcl-2 protein with an amino acid sequence comprising at least two
 CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
 CC shown in the specification. The invention also comprises an assay for
 CC identifying substances that bind to the Bcl-2 protein. The protein
 CC sequences of the invention are useful in biological assays to identify
 CC substances that block the ability of Bcl-2 to inhibit programmed cell
 CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.

XX Sequence 21 AA;

Query Match 100.0%; Score 83; DB 23; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.8e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRLRMSDFVD 16

|||||

Db 6 QRYGRLRMSDFVD 21

RESULT 6

AAU78630
 ID AAU78630 standard; Peptide: 21 AA.

XX AC AAU78630;

DT 18-JUN-2002 (first entry)

XX Human Bad peptide #30 which binds to a member of the Bcl-2 family.

XX Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
 KW ischemic injury; suppressor; BH3 domain.

XX Homo sapiens.

XX PN WO200220568-A2.
XX
XX PD 14-MAR-2002.
XX PF 04-SEP-2001; 2001WO-US27410.
XX PS 06-SEP-2000; 2000US-0656399.
XX PA (ABBO) ABBOTT LAB.
XX PI Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
XX PI Nettekheim DG, Swift KM, Matayoshi E, Zhang H;
XX DR WPI; 2002-292254/33.
XX
XX PT New derivatives of Bad peptide, useful for identifying compounds that
XX PT bind to Bcl-2 proteins, potential agents for treating cancer and
XX PT degenerative disease -
XX PS Claim 15; Page 18; 31pp; English.
XX
XX CC The present invention relates to new peptides that are derived from a
XX CC wild-type human Bad peptide and are able to bind to a member of the
XX CC Bcl-2 protein family. The peptides are useful, when labelled, in
XX CC competitive/displacement assays for identifying substances that bind to
XX CC members of the Bcl-2 family and may induce or suppress apoptosis so are
XX CC potentially useful for treating cancer (inducers) or degenerative
XX CC diseases or ischemic injury (suppressors). The peptides of the invention
XX CC have high helix propensity, maintain the contacts of the wild-type Bad
XX CC peptide and, compared with the Bad peptide, may have better physical
XX CC properties, particularly solubility. The present sequence represents one
XX CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
XX CC from the BH3 domain of the human wild-type Bad peptide.
XX
XX SQ Sequence 21 AA;
XX
XX Query Match 100.0%; Score 83; DB 23; Length 21;
XX Best Local Similarity 100.0%; Pred. No. 1.8e-07;
XX Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 QRYGRLRRMSDEFVD 16
XX Db 6 QRYGRLRRMSDEFVD 21
XX
XX RESULT 7
XX AAU78629
XX ID AAU78629 standard; Peptide; 22 AA.
XX AC
XX XX
XX DT 18-JUN-2002 (first entry)
XX DE Human Bad peptide #29 which binds to a member of the Bcl-2 family.
XX XX
XX KW Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
XX KW ischemic injury; suppressor; BH3 domain.
XX XX Homo sapiens.
XX OS
XX PN WO200220568-A2.
XX PD 14-MAR-2002.
XX PF 04-SEP-2001; 2001WO-US27410.
XX PS 06-SEP-2000; 2000US-0656399.
XX PA (ABBO) ABBOTT LAB.
XX PI Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
XX PI Nettekheim DG, Swift KM, Matayoshi E, Zhang H;
XX
XX

XX DR WPI; 2002-292254/33.
XX
XX PT New derivatives of Bad peptide, useful for identifying compounds that
XX PT bind to Bcl-2 proteins, potential agents for treating cancer and
XX PT degenerative disease -
XX PS Claim 15; Page 18; 31pp; English.
XX
XX CC The present invention relates to new peptides that are derived from a
XX CC wild-type human Bad peptide and are able to bind to a member of the
XX CC Bcl-2 protein family. The peptides are useful, when labelled, in
XX CC competitive/displacement assays for identifying substances that bind to
XX CC members of the Bcl-2 family and may induce or suppress apoptosis so are
XX CC potentially useful for treating cancer (inducers) or degenerative
XX CC diseases or ischemic injury (suppressors). The peptides of the invention
XX CC have high helix propensity, maintain the contacts of the wild-type Bad
XX CC peptide and, compared with the Bad peptide, may have better physical
XX CC properties, particularly solubility. The present sequence represents one
XX CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
XX CC from the BH3 domain of the human wild-type Bad peptide.
XX
XX SQ Sequence 22 AA;
XX
XX Query Match 100.0%; Score 83; DB 23; Length 22;
XX Best Local Similarity 100.0%; Pred. No. 1.9e-07;
XX Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 QRYGRLRRMSDEFVD 16
XX Db 6 QRYGRLRRMSDEFVD 21
XX
XX RESULT 8
XX AAU78628
XX ID AAU78628 standard; Peptide; 23 AA.
XX AC
XX XX
XX DT 18-JUN-2002 (first entry)
XX DE Human Bad peptide #28 which binds to a member of the Bcl-2 family.
XX XX
XX KW Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
XX KW ischemic injury; suppressor; BH3 domain.
XX XX Homo sapiens.
XX OS
XX PN WO200220568-A2.
XX PD 14-MAR-2002.
XX PF 04-SEP-2001; 2001WO-US27410.
XX PS 06-SEP-2000; 2000US-0656399.
XX PA (ABBO) ABBOTT LAB.
XX PI Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
XX PI Nettekheim DG, Swift KM, Matayoshi E, Zhang H;
XX DR WPI; 2002-292254/33.
XX
XX PT New derivatives of Bad peptide, useful for identifying compounds that
XX PT bind to Bcl-2 proteins, potential agents for treating cancer and
XX PT degenerative disease -
XX PS Claim 15; Page 18; 31pp; English.
XX
XX CC The present invention relates to new peptides that are derived from a
XX CC wild-type human Bad peptide and are able to bind to a member of the
XX CC Bcl-2 protein family. The peptides are useful, when labelled, in
XX CC competitive/displacement assays for identifying substances that bind to
XX CC members of the Bcl-2 family and may induce or suppress apoptosis so are
XX CC potentially useful for treating cancer (inducers) or degenerative
XX CC diseases or ischemic injury (suppressors). The peptides of the invention
XX CC have high helix propensity, maintain the contacts of the wild-type Bad
XX CC peptide and, compared with the Bad peptide, may have better physical
XX CC properties, particularly solubility. The present sequence represents one
XX CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
XX CC from the BH3 domain of the human wild-type Bad peptide.
XX
XX

CC members of the Bcl-2 family and may induce or suppress apoptosis so are
 CC potentially useful for treating cancer (inducers) or degenerative
 CC diseases of ischemic injury (suppressors). The peptides of the invention
 CC have high helix propensity, maintain the contacts of the wild-type Bad
 CC peptide and, compared with the Bad peptide, may have better physical
 CC properties, particularly solubility. The present sequence represents one
 CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
 CC from the BH3 domain of the human wild-type Bad peptide.

XX Sequence 23 AA;

SQ Query Match 100.0%; Score 83; DB 23; Length 23;
 Best Local Similarity 100.0%; Pred. No. 1.9e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDEFVD 16
 |||||
 Db 6 QRYGRELRRMSDEFVD 21

RESULT 9

ID ABG78482 standard; Peptide: 24 AA.

AC ABG78482;

DT 15-NOV-2002 (first entry)

XX Human Bcl2 fluorescein labelled peptide #2.

DE Human; Bcl2; BclX1; programmed cell death; apoptosis; mutant; muteln.

KW Homo sapiens.

OS Synthetic.

XX WO200240530-A2.

PN 23-MAY-2002.

XX 15-NOV-2001; 2001WO-US45693.

PF 20-NOV-2000; 2000US-0716395.

XX (ABBO) ABBOTT LAB.

PA Pesik SW, Petros AM, Yoon H, Nettesheim DG;

XX WPI: 2002-490141/52.

XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
 PT useful in biological assays to identify substances that block the
 PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -

XX Example 2; Page 15; 36pp; English.

XX This invention relates to a novel mutant protein which is derived from
 CC a wild type human Bcl-2 protein. The mutant is created by replacing a
 CC sequence of amino acid residues comprising a flexible loop from the wild
 CC type Bcl-2 protein with an amino acid sequence comprising at least two
 CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
 CC shown in the specification. The invention also comprises an assay for
 CC identifying substances that bind to the Bcl-2 protein. The protein
 CC sequences of the invention are useful in biological assays to identify
 CC substances that block the ability of Bcl-2 to inhibit programmed cell
 CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.

XX Sequence 24 AA;

SQ Query Match 100.0%; Score 83; DB 23; Length 24;
 Best Local Similarity 100.0%; Pred. No. 2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDEFVD 16
 |||||
 Db 6 QRYGRELRRMSDEFVD 21

RESULT 10

AAU78605
 ID AAU78605 standard; Peptide: 24 AA.

XX AAU78605;

DT 18-JUN-2002 (first entry)

XX Human Bad peptide #5 which binds to a member of the Bcl-2 family.

DE Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;

KW ischemic injury; suppressor; BH3 domain.

XX Homo sapiens.

XX Key Location/Qualifiers

EH Misc-difference 1
 FT /note= "Optionally labelled with 5-FAM,
 FT (6-carboxy-fluorescein)"

XX WO200220568-A2.

XX 14-MAR-2002.

XX 04-SEP-2001; 2001WO-US27410.

XX 06-SEP-2000; 2000US-0656399.

XX (ABBO) ABBOTT LAB.

XX Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;

PI Nettesheim DG, Swift KM, Matayoshi E, Zhang H;

XX WPI: 2002-292254/33.

XX New derivatives of Bad peptide, useful for identifying compounds that
 PT bind to Bcl-2 proteins, potential agents for treating cancer and
 PT degenerative disease -

XX Claim 4; Page 16; 31pp; English.

XX The present invention relates to new peptides that are derived from a
 CC wild-type human Bad peptide and are able to bind to a member of the
 CC Bcl-2 protein family. The peptides are useful, when labelled, in
 CC competitive/displacement assays for identifying substances that bind to
 CC members of the Bcl-2 family and may induce or suppress apoptosis so are
 CC potentially useful for treating cancer (inducers) or degenerative
 CC diseases or ischemic injury (suppressors). The peptides of the invention
 CC have high helix propensity, maintain the contacts of the wild-type Bad
 CC peptide and, compared with the Bad peptide, may have better physical
 CC properties, particularly solubility. The present sequence represents one
 CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
 CC from the BH3 domain of the human wild-type Bad peptide.

XX Sequence 24 AA;

SQ Query Match 100.0%; Score 83; DB 23; Length 24;
 Best Local Similarity 100.0%; Pred. No. 2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDEFVD 16
 |||||
 Db 6 QRYGRELRRMSDEFVD 21

RESULT 11

AAU78627
 ID AAU78627 standard; Peptide: 24 AA.

XX AC AAU78627;
 XX DT 18-JUN-2002 (first entry)
 XX DE Human Bad peptide #27 which binds to a member of the Bcl-2 family.
 XX DE Human; Bad; Bcl-2; apoptosis; cancer; inducer; degenerative disease;
 KW ischaemic injury; suppressor; BH3 domain.
 XX OS Homo sapiens.
 XX PN WO200220568-A2.
 XX PD 14-MAR-2002.
 XX PF 04-SEP-2001; 2001WO-US27410.
 XX PR 06-SEP-2000; 2000US-0656399.
 XX PA (ABBO) ABBOTT LAB.
 XX PI Fesik SW, Meadows RP, Joseph MK, Olejniczak ET, Petros AM;
 PI Nettesheim DG, Swift KM, Matayoshi E, Zhang H;
 XX DR WPI; 2002-292254/33.
 XX DE New derivatives of Bad peptide, useful for identifying compounds that
 PT bind to Bcl-2 proteins, potential agents for treating cancer and
 PT degenerative disease -
 XX PS Claim 15; Page 18; 31pp; English.
 XX CC The present invention relates to new peptides that are derived from a
 CC wild-type human Bad peptide and are able to bind to a member of the
 CC Bcl-2 protein family. The peptides are useful, when labelled, in
 CC competitive/displacement assays for identifying substances that bind to
 CC members of the Bcl-2 family and may induce or suppress apoptosis so are
 CC potentially useful for treating cancer (inducers) or degenerative
 CC diseases or ischaemic injury (suppressors). The peptides of the invention
 CC have high helix propensity, maintain the contacts of the wild-type Bad
 CC peptide and, compared with the Bad peptide, may have better physical
 CC properties, particularly solubility. The present sequence represents one
 CC of a collection of Bad peptides (AAU78601-AAU78631) that were derived
 CC from the BH3 domain of the human wild-type Bad peptide.
 XX SQ Sequence 24 AA;
 Query Match 100.0%; Score 83; DB 23; Length 24;
 Best Local Similarity 100.0%; Pred. No. 2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRLRRMSDEFVD 16
 DB 6 QRYGRLRRMSDEFVD 21
 RESULT 12
 ABP56161
 ID ABP56161 standard; peptide; 25 AA.
 XX AC ABP56161;
 XX DT 28-MAR-2003 (first entry)
 XX DE PTPC-interacting TOX peptide #27.
 KW Mitochondrial membrane permeabilisation; mitochondrion; PTPC;
 KW permeability transition pore complex; virucide; neuroprotective;
 KW vasotrophic; cytostatic; infection; cell death regulation; apoptosis;
 KW mitochondrial permeability transition pore complex modulator; cancer;
 KW apoptogenic; ischaemia; neurodegenerative disease; fulminant hepatitis.
 XX

OS Synthetic.
 XX PN WO200261105-A2.
 XX PD 08-AUG-2002.
 XX PF 01-FEB-2002; 2002WO-EP01633.
 XX PR 02-FEB-2001; 2001US-265594P.
 XX PA (INSP) INST PASTEUR.
 PA (CNRS) CENT NAT RECH SCI.
 XX PI Edelman L, Jacotot E, Briand J;
 XX DR WPI; 2002-619260/66.
 XX DE New chimeric bifunctional molecules that target specific cells and
 PT regulate the apoptosis function of the permeability transition pore
 PT complex of the mitochondria, useful for treating or preventing e.g.
 PT cancer or ischemia -
 XX PS Claim 9; Page 11; 76pp; English.
 XX CC The present invention describes a chimeric bifunctional molecule (I)
 CC comprising at least a first functional molecule covalently linked to a
 CC second functional molecule, which is able to modulate the activity of
 CC the permeability transition pore complex (PTPC) of the mitochondria.
 CC (I) has the function of specifically targeting and entering a tissue
 CC cell population. The second functional molecule has the function of
 CC specifically targeting, and inducing or preventing the death of the
 CC cells by apoptosis by regulating the opening or the closing of the PTPC
 CC of the mitochondria or its fragment. (I) has virucide, neuroprotective,
 CC vasotropic and cytostatic activities, and can be used as a mitochondrial
 CC permeability transition pore complex (PTPC) modulator. (I) is useful for
 CC treating or preventing a pathological infection or disease. (I) is also
 CC useful for regulating cell death regulatory molecules, specifically the
 CC apoptogenic function of the PTPC, for treating e.g. cancer, ischaemia,
 CC neurodegenerative diseases, fulminant hepatitis or viral infections.
 CC The present sequence represents a PTPC-interacting TOX peptide which is
 CC given in the exemplification of the present invention.
 XX SQ Sequence 25 AA;
 Query Match 100.0%; Score 83; DB 23; Length 25;
 Best Local Similarity 100.0%; Pred. No. 2.1e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRLRRMSDEFVD 16
 DB 6 QRYGRLRRMSDEFVD 21
 RESULT 13
 ABG78481
 ID ABG78481 standard; Peptide; 25 AA.
 XX AC ABG78481;
 XX DT 15-NOV-2002 (first entry)
 XX DE Human Bcl2 fluorescein labelled peptide #1.
 KW Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; muten.
 XX OS Homo sapiens.
 OS Synthetic.
 XX PN WO200240530-A2.
 XX PD 23-MAY-2002.
 XX PF 15-NOV-2001; 2001WO-US45693.

XX 20-NOV-2000; 2000US-0716395.
 XX (ABBO) ABBOTT LAB.
 XX Fesik SW, Petros AM, Yoon H, Nettesheim DG;
 XX WPI; 2002-490141/52.
 XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
 PT useful in biological assays to identify substances that block the
 PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -
 XX
 XX Example 2; Page 15; 36pp; English.
 XX This invention relates to a novel mutant protein which is derived from
 CC a wild type human Bcl-2 protein. The mutant is created by replacing a
 CC sequence of amino acid residues comprising a flexible loop from the wild
 CC type Bcl-2 protein with an amino acid sequence comprising at least two
 CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
 CC shown in the specification. The invention also comprises an assay for
 CC identifying substances that bind to the Bcl-2 protein. The protein
 CC sequences of the invention are useful in biological assays to identify
 CC substances that block the ability of Bcl-2 to inhibit programmed cell
 CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.
 XX
 XX Sequence 25 AA;
 XX
 XX Query Match 100.0%; Score 83; DB 23; Length 25;
 XX Best Local Similarity 100.0%; Pred. No. 2.1e-07;
 XX Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRRMSDEVD 16
 DB 6 QRYGRELRRMSDEVD 21
 RESULT 14
 ABG78484
 ID ABG78484 standard; Peptide; 25 AA.
 XX
 XX ABG78484;
 XX 15-NOV-2002 (first entry)
 XX Mutant Bcl2 competitive binding assay peptide #1.
 XX Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; muteln.
 XX Homo sapiens.
 XX Synthetic.
 XX W0200240530-A2.
 XX 23-MAY-2002.
 XX 15-NOV-2001; 2001WO-US45693.
 XX 20-NOV-2000; 2000US-0716395.
 XX (ABBO) ABBOTT LAB.
 XX Fesik SW, Petros AM, Yoon H, Nettesheim DG;
 XX WPI; 2002-490141/52.
 XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
 PT useful in biological assays to identify substances that block the
 PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -
 XX
 XX Example 2; Page 17; 36pp; English.

CC This invention relates to a novel mutant protein which is derived from
 CC a wild type human Bcl-2 protein. The mutant is created by replacing a
 CC sequence of amino acid residues comprising a flexible loop from the wild
 CC type Bcl-2 protein with an amino acid sequence comprising at least two
 CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
 CC shown in the specification. The invention also comprises an assay for
 CC identifying substances that bind to the Bcl-2 protein. The protein
 CC sequences of the invention are useful in biological assays to identify
 CC substances that block the ability of Bcl-2 to inhibit programmed cell
 CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.
 XX
 XX Sequence 25 AA;
 XX
 XX Query Match 100.0%; Score 83; DB 23; Length 25;
 XX Best Local Similarity 100.0%; Pred. No. 2.1e-07;
 XX Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRELRRMSDEVD 16
 DB 6 QRYGRELRRMSDEVD 21
 RESULT 15
 ABG78485
 ID ABG78485 standard; Peptide; 25 AA.
 XX
 XX ABG78485;
 XX 15-NOV-2002 (first entry)
 XX Mutant Bcl2 competitive binding assay peptide #2.
 XX Human; Bcl2; BclXl; programmed cell death; apoptosis; mutant; muteln.
 XX Homo sapiens.
 XX Synthetic.
 XX W0200240530-A2.
 XX 23-MAY-2002.
 XX 15-NOV-2001; 2001WO-US45693.
 XX 20-NOV-2000; 2000US-0716395.
 XX (ABBO) ABBOTT LAB.
 XX Fesik SW, Petros AM, Yoon H, Nettesheim DG;
 XX WPI; 2002-490141/52.
 XX New mutant Bcl-2 proteins derived from a wild type human Bcl-2 protein,
 PT useful in biological assays to identify substances that block the
 PT ability of Bcl-2 to inhibit programmed cell death or apoptosis -
 XX
 XX Example 2; Page 17; 36pp; English.
 XX This invention relates to a novel mutant protein which is derived from
 CC a wild type human Bcl-2 protein. The mutant is created by replacing a
 CC sequence of amino acid residues comprising a flexible loop from the wild
 CC type Bcl-2 protein with an amino acid sequence comprising at least two
 CC acidic amino acids. The mutant Bcl-2 protein comprises a 166 residue
 CC shown in the specification. The invention also comprises an assay for
 CC identifying substances that bind to the Bcl-2 protein. The protein
 CC sequences of the invention are useful in biological assays to identify
 CC substances that block the ability of Bcl-2 to inhibit programmed cell
 CC death or apoptosis. The present sequence represents a human Bcl2
 CC peptide of the invention.
 XX
 XX Sequence 25 AA;
 XX
 XX Query Match 100.0%; Score 83; DB 23; Length 25;

Best Local Similarity 100.0%; Pred. No. 2.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRELRRMSDEFVD 16
 |||||
Db 6 QRYGRELRRMSDEFVD 21

Search completed: September 15, 2003, 17:22:14
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(without alignments)
81.144 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRLRMSDFVD 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents.AA.*

- 1: /cgn2_6/prodata/1/iaa/5A.COMB.pep.*
- 2: /cgn2_6/prodata/1/iaa/5B.COMB.pep.*
- 3: /cgn2_6/prodata/1/iaa/6A.COMB.pep.*
- 4: /cgn2_6/prodata/1/iaa/6B.COMB.pep.*
- 5: /cgn2_6/prodata/1/iaa/PCTUS.COMB.pep.*
- 6: /cgn2_6/prodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	83	100.0	166	1	US-08-665-617-2
2	83	100.0	168	1	Sequence 2, Appli
3	83	100.0	168	3	Sequence 1, Appli
4	83	100.0	168	3	Sequence 7, Appli
5	83	100.0	168	3	Sequence 1, Appli
6	83	100.0	168	3	Sequence 7, Appli
7	83	100.0	168	4	US-09-375-257-2
8	73	88.0	23	1	US-08-333-565-10
9	73	88.0	23	2	US-08-661-479-10
10	73	88.0	59	2	US-08-733-505A-55
11	73	88.0	59	2	US-08-733-505A-56
12	73	88.0	59	2	US-08-733-505A-57
13	73	88.0	59	2	US-08-733-505A-58
14	73	88.0	204	1	US-08-333-565-2
15	73	88.0	204	1	US-08-661-479-2
16	73	88.0	204	2	US-08-733-505A-1
17	73	88.0	204	2	US-08-733-505A-12
18	73	88.0	204	2	US-08-733-505A-13
19	73	88.0	204	2	US-08-733-505A-14
20	70	84.3	204	2	US-08-717-123-3
21	70	84.3	204	4	US-09-375-257-3
22	67	80.7	16	1	US-08-333-565-26
23	67	80.7	16	2	US-08-661-479-26
24	44	53.0	1125	4	US-09-252-991A-18729
25	42	50.6	11	2	US-08-733-505A-34
26	42	50.6	11	2	US-08-706-741B-69
27	42	50.6	11	2	US-08-924-695A-69

28 42 50.6 876 1 US-08-785-429-2 Sequence 2, Appli
29 42 50.6 876 3 US-08-996-621-2 Sequence 2, Appli
30 42 50.6 888 4 US-09-134-001C-3032 Sequence 3032, Ap
31 41 49.4 284 4 US-08-328-352-6559 Sequence 6559, Ap
32 40 48.2 394 4 US-09-252-991A-31592 Sequence 31592, A
33 40 48.2 724 4 US-08-328-352-7710 Sequence 7710, Ap
34 39 47.0 333 4 US-09-252-991A-28443 Sequence 28443, A
35 38.5 46.4 1112 4 US-09-252-991A-27256 Sequence 27256, A
36 38 45.8 376 4 US-09-252-991A-26270 Sequence 26270, A
37 38 45.8 432 3 US-09-075-087-2 Sequence 2, Appli
38 38 45.8 432 3 US-09-472-371-1 Sequence 1, Appli
39 38 45.8 575 3 US-08-913-805A-2 Sequence 2, Appli
40 38 45.8 575 3 US-08-913-805A-10 Sequence 10, Appli
41 38 45.8 575 3 US-09-442-629-2 Sequence 2, Appli
42 38 45.8 575 3 US-09-442-629-10 Sequence 10, Appli
43 37 44.6 338 4 US-09-252-991A-30165 Sequence 30165, A
44 37 44.6 520 4 US-09-391-104-10 Sequence 10, Appli
45 36 43.4 80 4 US-09-328-352-8004 Sequence 8004, Ap

ALIGNMENTS

RESULT 1
US-08-665-617-2
: Sequence 2, Application US/08665617
: Patent No. 5663316
: GENERAL INFORMATION:
: APPLICANT: Xudong, Yin
: TITLE OF INVENTION: Gene and Protein for Regulation of Cell Death
: NUMBER OF SEQUENCES: 2
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Saliwanchik & Saliwanchik
: STREET: 2421 N.W. 41st Street, Suite A-1
: CITY: Gainesville
: STATE: Florida
: COUNTRY: USA
: ZIP: 32606
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/665,617
: FILING DATE:
: CLASSIFICATION: 530
: ATTORNEY/AGENT INFORMATION:
: NAME: Saliwanchik, David R.
: REGISTRATION NUMBER: 31,794
: REFERENCE/DOCKET NUMBER: CL-8
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (352) 375-8100
: TELEFAX: (352) 372-5800
: INFORMATION FOR SEQ ID NO: 2:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 166 amino acids
: TYPE: amino acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: protein
US-08-665-617-2

Query Match 100.0%; Score 83; DB 1; Length 166;
Best local similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QRYGRLRMSDFVD 16

Db 106 QRYGRLRMSDFVD 121

RESULT 2

```
US-08-717-123-2
; Sequence 2, Application US/08/17123
; Patent No. 5965703
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: Human BAD Polypeptides, Encoding Nucleic
; TITLE OF INVENTION: Acids and Methods of Use
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/717,123
; FILING DATE: 20-SEP-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-ID 1929
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
;
US-08-717-123-2
Query Match 100.0%; Score 83; DB 2; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEVD 16
| | | | | | | | | | | | | | | |
Db 108 QRYGRELRRMSDEVD 123

RESULT 3
US-08-985-335-1
; Sequence 1, Application US/08/985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 388673
;
US-08-985-335-1
Query Match 100.0%; Score 83; DB 3; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEVD 16
| | | | | | | | | | | | | | | |
Db 108 QRYGRELRRMSDEVD 123

RESULT 4
US-08-985-335-7
; Sequence 7, Application US/08/985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
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; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 388673
;
US-08-985-335-1
Query Match 100.0%; Score 83; DB 3; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEVD 16
| | | | | | | | | | | | | | | |
Db 108 QRYGRELRRMSDEVD 123

RESULT 4
US-08-985-335-7
; Sequence 7, Application US/08/985335
; Patent No. 6080847
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Lal, Preeti
; APPLICANT: Shah, Purvi
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; TITLE OF INVENTION: PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,335
; FILING DATE: Filed Herewith
; PRIOR APPLICATION NUMBER:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
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SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 1683637
US-08-985-335-7

Query Match 100.0%; Score 83; DB 3; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFVD 16
|||||

Db 108 QRYGRELRRMSDEFVD 123

RESULT 5

US-09-410-372-1
Sequence 1, Application US/09410372
Patent No. 6281334
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
TITLE OF INVENTION: PROLIFERATION
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/410,372
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/985,335
FILING DATE:

ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0421 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166

INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: SYNORAB01
CLONE: 358673
US-09-410-372-1

Query Match 100.0%; Score 83; DB 3; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFVD 16

Db 108 QRYGRELRRMSDEFVD 123
|||||

RESULT 6

US-09-410-372-7
Sequence 7, Application US/09410372
Patent No. 6281334
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
TITLE OF INVENTION: PROLIFERATION
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/410,372
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/985,335
FILING DATE:

ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0421 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166

INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 168 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 1683637
US-09-410-372-7

Query Match 100.0%; Score 83; DB 3; Length 168;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFVD 16
|||||

Db 108 QRYGRELRRMSDEFVD 123

RESULT 7

US-09-375-257-2
Sequence 2, Application US/09375257
Patent No. 6504022
GENERAL INFORMATION:
APPLICANT: Horne, William A.
APPLICANT: Oltersdorf, Tilman
TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
TITLE OF INVENTION: ACIDS AND METHODS OF USE
FILE REFERENCE: 480140.428D1
CURRENT APPLICATION NUMBER: US/09/375,257

; CURRENT FILING DATE: 1999-08-16
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-375-257-2

Query Match 100.0%; Score 83; DB 4; Length 168;
Best Local Similarity 100.0%; Pred. No. 12.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 16
 |||||
DB 108 QRYGRELRRMSDEF 123

RESULT 8
US-08-333-565-10
; Sequence 10, Application US/08333565
; Patent No. 5622852
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Kourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/333,565
; FILING DATE: 31-OCT-1994
; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 23 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-333-565-10

Query Match 88.0%; Score 73; DB 1; Length 23;
Best Local Similarity 100.0%; Pred. No. 1.5e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
 |||||
DB 8 QRYGRELRRMSDEF 21

RESULT 9
US-08-661-479-10
; Sequence 10, Application US/08661479
; Patent No. 5834209
; GENERAL INFORMATION:

; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: Bcl-x/Bcl-2 ASSOCIATED CELL DEATH
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Kourie and Crew
; STREET: 379 Lytton Avenue
; CITY: Palo Alto
; STATE: California
; COUNTRY: US
; ZIP: 94301

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/661,479
; FILING DATE: 11-JUN-1995
; CLASSIFICATION: 435

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/333,565
; FILING DATE: 31-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 15726A-000700
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 326-2400
; TELEFAX: (415) 326-2422
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 23 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-661-479-10

Query Match 88.0%; Score 73; DB 2; Length 23;
Best Local Similarity 100.0%; Pred. No. 1.5e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
 |||||
DB 8 QRYGRELRRMSDEF 21

RESULT 10
US-08-733-505A-55
; Sequence 55, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, Stanley J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:
NAME: HOLLAND, DONALD R.
REGISTRATION NUMBER: 35,197
REFERENCE/DOCKET NUMBER: 965458
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314) 727-5188
TELEFAX: (314) 727-6092
INFORMATION FOR SEQ ID NO: 55:
SEQUENCE CHARACTERISTICS:
LENGTH: 59 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-733-505A-55

Query Match 88.0%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 4.2e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
|||||
Db 46 QRYGRELRRMSDEF 59

RESULT 11
US-08-733-505A-56
; Sequence 56, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 56:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-56

Query Match 88.0%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 4.2e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
|||||
Db 46 QRYGRELRRMSDEF 59

RESULT 12
US-08-733-505A-57
; Sequence 57, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/733,505A
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 965458
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 57:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-733-505A-57

Query Match 88.0%; Score 73; DB 2; Length 59;
Best Local Similarity 100.0%; Pred. No. 4.2e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
|||||
Db 46 QRYGRELRRMSDEF 59

RESULT 13
US-08-733-505A-58
; Sequence 58, Application US/08733505A
; Patent No. 5856445
; GENERAL INFORMATION:
; APPLICANT: KORSMEYER, STANLEY J.
; TITLE OF INVENTION: SERINE SUBSTITUTED MUTANTS OF
; TITLE OF INVENTION: BCL-XL/BCL-2 ASSOCIATED CELL DEATH REGULATOR
; NUMBER OF SEQUENCES: 60
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BLVD., SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: USA
; ZIP: 63105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30


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TELEPHONE: (415) 326-2400
TELEFAX: (415) 326-2422
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 204 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
FEATURE:
NAME/KEY: Protein
LOCATION: 1..204
OTHER INFORMATION: /note= "Deduced amino
OTHER INFORMATION: of mouse BAD."

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Search completed: September 15, 2003, 17:45:06
Job time : 8.34286 secs

genCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 15, 2003, 17:25:56 ; Search time 12.6857 Seconds
(without alignments)
184.034 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRELRRMSDFVD 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 541936 seqs, 145912426 residues

Total number of hits satisfying chosen parameters: 541936

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 10%

Listing first 45 summaries

Database : Published Applications AA:*

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14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Score	Match	Length	DB ID	Description
1	83	100.0	25	15	US-10-059-261-258
2	83	100.0	168	9	US-09-922-378-2
3	83	100.0	168	9	US-09-894-657-1
4	83	100.0	168	9	US-09-894-657-7
5	83	100.0	168	14	US-10-066-179-2
6	70	84.3	204	9	US-09-922-378-3
7	70	84.3	204	14	US-10-066-179-3
8	56	67.5	15	15	US-10-174-105A-147
9	44	53.0	215	15	US-10-156-761-9145
10	42	50.6	647	15	US-10-156-761-9145
11	42	50.6	810	9	US-09-815-242-12636
12	42	50.6	876	9	US-09-815-242-12636
13	42	50.6	879	9	US-09-815-242-13003
14	41	49.4	312	15	US-10-156-761-7630
15	41	49.4	312	15	US-10-156-761-7867

16 40.5 48.8 1265 15 US-10-138-070-69 Sequence 69, Appl
17 40 48.2 113 9 US-09-815-242-13424 Sequence 13424, A
18 40 48.2 380 15 US-10-156-761-13594 Sequence 13594, A
19 39 47.0 454 15 US-10-166-087-46 Sequence 46, Appl
20 39 47.0 959 12 US-10-342-224-40 Sequence 40, Appl
21 38.5 46.4 588 15 US-10-156-761-10330 Sequence 10330, A
22 38 45.8 35 15 US-10-032-750-1 Sequence 1, Appl
23 38 45.8 50 10 US-09-971-980-64 Sequence 64, Appl
24 38 45.8 64 10 US-09-971-980-62 Sequence 62, Appl
25 38 45.8 138 15 US-10-092-750-241 Sequence 241, App
26 38 45.8 313 10 US-09-738-626-3724 Sequence 3724, Ap
27 38 45.8 320 10 US-09-738-626-4522 Sequence 4522, Ap
28 38 45.8 515 15 US-10-106-698-4658 Sequence 4658, Ap
29 38 45.8 575 9 US-09-839-136-2 Sequence 2, Appl
30 38 45.8 575 9 US-09-839-136-10 Sequence 10, Appl
31 37 44.6 391 15 US-10-163-198-19 Sequence 19, Appl
32 37 44.6 543 15 US-10-156-761-9820 Sequence 9820, Ap
33 37 44.6 556 12 US-09-949-029-36 Sequence 36, Appl
34 37 44.6 1270 15 US-10-223-070-8 Sequence 8, Appl
35 36 43.4 78 15 US-10-083-357-1237 Sequence 1237, Ap
36 36 43.4 80 15 US-10-106-698-5539 Sequence 5539, Ap
37 36 43.4 103 10 US-09-738-626-4453 Sequence 4453, Ap
38 36 43.4 270 11 US-09-934-455-162 Sequence 162, App
39 36 43.4 284 15 US-10-156-761-10335 Sequence 10335, A
40 36 43.4 292 10 US-09-738-626-4426 Sequence 4426, Ap
41 36 43.4 350 11 US-09-940-244-394 Sequence 394, App
42 36 43.4 393 10 US-09-950-510-22 Sequence 22, Appl
43 36 43.4 394 10 US-09-712-363-205 Sequence 205, App
44 36 43.4 404 15 US-10-156-761-14659 Sequence 14659, A
45 36 43.4 445 10 US-09-950-510-12 Sequence 12, Appl

ALIGNMENTS

RESULT 1

; Sequence 258, Application US/10059261
; Publication No. US20030077826A1
; GENERAL INFORMATION:
; APPLICANT: EDELMAN, LENA
; APPLICANT: JACOTOT, ETIENNE DANIEL FRANCOIS
; APPLICANT: BRIAND, JEAN-PAUL
; TITLE OF INVENTION: CHIMERIC MOLECULES CONTAINING A MODULE ABLE TO TARGET
; TITLE OF INVENTION: SPECIFIC CELLS AND A MODULE REGULATING THE APOPTOTIC
; TITLE OF INVENTION: FUNCTION OF THE PERMEABILITY TRANSITION PORE COMPLEX
; TITLE OF INVENTION: (PTPC)
; FILE REFERENCE: 03495.0216
; CURRENT APPLICATION NUMBER: US/10/059,261
; CURRENT FILING DATE: 2002-08-29
; PRIOR APPLICATION NUMBER: 60/265,594
; PRIOR FILING DATE: 2001-02-02
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 258
; LENGTH: 25
; TYPE: PRT
; ORGANISM: Unknown Organism
; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: TOX peptide
US-10-059-261-258

Query Match 100.0%; Score 83; DB 15; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDFVD 16

|||||

Db 6 QRYGRELRRMSDFVD 21

RESULT 2

US-09-922-378-2

```
; Sequence 2, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN RAD POLYPPPTIDES, ENCODING NUCLEIC
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-922-378-2

Query Match      100.0%; Score 83; DB 9; Length 168;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRELRLMSDEVD 16
Db      108 QRYGRELRLMSDEVD 123

RESULT 3
US-09-894-657-1
; Sequence 1, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: SYNORAB01
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-894-657-1

Query Match      100.0%; Score 83; DB 9; Length 168;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRELRLMSDEVD 16
Db      108 QRYGRELRLMSDEVD 123

RESULT 4
US-09-894-657-7
; Sequence 7, Application US/09894657
; Patent No. US20020098569A1
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; Yue, Henry
; Lal, Preeti
; Shah, Purvi
; Corley, Neil C.
; TITLE OF INVENTION: PROTEINS ASSOCIATED WITH CELL
; PROLIFERATION
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Dr.
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/894,657
; FILING DATE: 28-Jun-2001
; PRIOR APPLICATION DATA:
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0421 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 168 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 1683637
; SEQUENCE DESCRIPTION: SEQ ID NO: 7:
US-09-894-657-7

Query Match      100.0%; Score 83; DB 9; Length 168;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 QRYGRELRLMSDEVD 16
Db      108 QRYGRELRLMSDEVD 123

RESULT 5
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US-10-066-179-2
; Sequence 2, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 168
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-066-179-2

Query Match      100.0%; Score 83; DB 14; Length 168;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFVD 16
   |||||
Db 108 QRYGRELRRMSDEFVD 123

RESULT 6
US-09-922-378-3
; Sequence 3, Application US/09922378
; Patent No. US20020037869A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428D3
; CURRENT APPLICATION NUMBER: US/09/922,378
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-922-378-3

Query Match      84.3%; Score 70; DB 9; Length 204;
Best Local Similarity 92.9%; Pred. No. 0.00064;
Matches 13; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
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Db 145 QRYGRELRRMTDEF 158

RESULT 7
US-10-066-179-3
; Sequence 3, Application US/10066179
; Publication No. US20020115631A1
; GENERAL INFORMATION:
; APPLICANT: Horne, William A.
; APPLICANT: Oltersdorf, Tilman
; TITLE OF INVENTION: HUMAN BAD POLYPEPTIDES, ENCODING NUCLEIC
; TITLE OF INVENTION: ACIDS AND METHODS OF USE
; FILE REFERENCE: 480140.428C1
; CURRENT APPLICATION NUMBER: US/10/066,179
; CURRENT FILING DATE: 2002-02-01
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 204
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-066-179-3

Query Match      84.3%; Score 70; DB 14; Length 204;
Best Local Similarity 92.9%; Pred. No. 0.00064;
Matches 13; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
   |||||
Db 145 QRYGRELRRMTDEF 158

RESULT 8
US-10-174-105A-147
; Sequence 147, Application US/10174105A
; Publication No. US20030068652A1
; GENERAL INFORMATION:
; APPLICANT: Cell Signaling Technology, Inc.
; APPLICANT: ZHANG, Hui
; APPLICANT: COMB, Michael J.
; APPLICANT: TAN, Yi
; TITLE OF INVENTION: POSITIVE IDENTIFICATION OF PHOSPHO-PROTEINS USING MOTIF-SPECI
; TITLE OF INVENTION: CONTEXT-INDEPENDENT ANTIBODIES COUPLED WITH DATABASE SEARCHI
; FILE REFERENCE: CST-138 CIP3
; CURRENT APPLICATION NUMBER: US/10/174,105A
; CURRENT FILING DATE: 2002-06-18
; PRIOR APPLICATION NUMBER: US 09/148,712
; PRIOR FILING DATE: 1998-09-04
; PRIOR APPLICATION NUMBER: US 09/535,364
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 193
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 147
; LENGTH: 15
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide
; NAME/KEY: MOD_RES
; LOCATION: (8)..(8)
; OTHER INFORMATION: PHOSPHORYLATION; serine at position 8 is phosphorylated
US-10-174-105A-147

Query Match      67.5%; Score 56; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.0084;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GRELRRMSDEF 14
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Db 1 GRELRRMSDEF 11

RESULT 9
US-10-156-761-9145
; Sequence 9145, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
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; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 9145
; LENGTH: 215
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-9145
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Query Match          53.0%; Score 44; DB 15; Length 215;
Best Local Similarity 61.5%; Pred. No. 12;
Matches      8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
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QY      1 QRYGRLRRMSDE 13
          :|:|:|:|:|:|
Db      108 ERWGGDLRRARDE 120
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```
RESULT 10
US-10-156-761-9735
; Sequence 9735, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 9735
; LENGTH: 647
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-9735
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Query Match          50.6%; Score 42; DB 15; Length 647;
Best Local Similarity 57.1%; Pred. No. 83;
Matches      8; Conservative 1; Mismatches 5; Indels 0; Gaps 0;
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QY      3 YGRLRRMSDEVD 16
          | | | | | | | |
Db      565 YARGLRHGHDELID 578
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RESULT 11
US-09-815-242-12636
; Sequence 12636, Application US/09815242
; Patent No. US20020061569A1
; GENERAL INFORMATION:
; APPLICANT: Haselbeck, Robert
; APPLICANT: Ohlsen, Kari L.
; APPLICANT: Zyskind, Judith W.
; APPLICANT: Wall, Daniel
; APPLICANT: Trawick, John D.
; APPLICANT: Carr, Grant J.
; APPLICANT: Yamamoto, Robert T.
; APPLICANT: Xu, H. Howard
; TITLE OF INVENTION: Identification of Essential Genes in
; FILE REFERENCE: ELITRA.011a
; CURRENT APPLICATION NUMBER: US/09/815,242
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: 60/191,078
; PRIOR FILING DATE: 2000-03-21
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; PRIOR APPLICATION NUMBER: 60/206,848
; PRIOR FILING DATE: 2000-05-23
; PRIOR APPLICATION NUMBER: 60/207,727
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: 60/242,578
; PRIOR FILING DATE: 2000-10-23
; PRIOR APPLICATION NUMBER: 60/253,625
; PRIOR FILING DATE: 2000-11-27
; PRIOR APPLICATION NUMBER: 60/257,931
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: 60/269,308
; PRIOR FILING DATE: 2001-02-16
; NUMBER OF SEQ ID NOS: 14110
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12636
; LENGTH: 810
; TYPE: PRT
; ORGANISM: Staphylococcus aureus
US-09-815-242-12636
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Query Match          50.6%; Score 42; DB 9; Length 810;
Best Local Similarity 61.5%; Pred. No. 11e+02;
Matches      8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
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QY      4 GRELRRMSDEVD 16
          | | | | | :|:|:|
Db      185 GRELPILADEYVD 197
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RESULT 12
US-09-815-242-13003
; Sequence 13003, Application US/09815242
; Patent No. US20020061569A1
; GENERAL INFORMATION:
; APPLICANT: Haselbeck, Robert
; APPLICANT: Ohlsen, Kari L.
; APPLICANT: Zyskind, Judith W.
; APPLICANT: Wall, Daniel
; APPLICANT: Trawick, John D.
; APPLICANT: Carr, Grant J.
; APPLICANT: Yamamoto, Robert T.
; APPLICANT: Xu, H. Howard
; TITLE OF INVENTION: Identification of Essential Genes in
; FILE REFERENCE: ELITRA.011a
; CURRENT APPLICATION NUMBER: US/09/815,242
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: 60/191,078
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: 60/206,848
; PRIOR FILING DATE: 2000-05-23
; PRIOR APPLICATION NUMBER: 60/207,727
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: 60/242,578
; PRIOR FILING DATE: 2000-10-23
; PRIOR APPLICATION NUMBER: 60/253,625
; PRIOR FILING DATE: 2000-11-27
; PRIOR APPLICATION NUMBER: 60/257,931
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: 60/269,308
; PRIOR FILING DATE: 2001-02-16
; NUMBER OF SEQ ID NOS: 14110
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13003
; LENGTH: 876
; TYPE: PRT
; ORGANISM: Staphylococcus aureus
US-09-815-242-13003
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Query Match          50.6%; Score 42; DB 9; Length 876;
Best Local Similarity 61.5%; Pred. No. 11e+02;
Matches      8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
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; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 7630
; LENGTH: 312
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-7630

Query Match 49.4%; Score 41; DB 15; Length 312;
Best Local Similarity 50.0%; Pred. No. 57;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDFVD 16
| | | | | : | : |
Db 125 QLSGEERRRLDDFMD 140

RESULT 15
US-10-156-761-7867
; Sequence 7867, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 7867
; LENGTH: 312
; TYPE: PRT
; ORGANISM: Streptomyces avermitilis
US-10-156-761-7867

Query Match 49.4%; Score 41; DB 15; Length 312;
Best Local Similarity 50.0%; Pred. No. 57;
Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDFVD 16
| | | | | : | : |
Db 125 QLSGEERRRLDDFMD 140

Search completed: September 15, 2003, 17:47:53
Job time : 13.6857 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:16:55 ; Search time 3.77143 Seconds
(without alignments)
199.507 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRELRRMSDFVD 16

Scoring table:

BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	83	100.0	168	1 BAD_HUMAN	Q92934 mus sapien
2	73	88.0	204	1 BAD_MOUSE	Q61337 mus musculu
3	73	88.0	205	1 BAD_RAT	Q35147 rattus norv
4	42	50.6	196	1 B1M_MOUSE	O54918 mus musculu
5	42	50.6	196	1 B1M_RAT	O88498 rattus norv
6	42	50.6	457	1 HGD_RHIL0	Q983j4 rhizobium l
7	42	50.6	653	1 HT2A_HUMAN	Q13049 homo sapien
8	41	49.4	370	1 AROG_CANAL	P79023 candida alb
9	41	49.4	861	1 GCR3_YEAST	P34160 saccharomyc
10	40	48.2	261	1 YGFG_ECOLI	P52045 escherichia
11	40	48.2	380	1 PHLC_TRYCR	O15886 trypanosoma
12	40	48.2	429	1 MTAL_ACSPA	O52702 acetobacter
13	40	48.2	631	1 RPSD_BORBU	P52323 borrelia bu
14	39	47.0	220	1 GPGL_THEMA	Q9x0n8 thermotoga
15	39	47.0	365	1 R4S1_SCHPO	P36601 schizosacch
16	39	47.0	418	1 V8I2_REOVD	P03525 reovirus (t
17	39	47.0	418	1 V8I2_REOVL	P11314 reovirus (t
18	39	47.0	432	1 TIG_SALTY	O8x1c4 salmonella
19	39	47.0	503	1 MTRB_BPRH1	P09915 bacterioph
20	39	47.0	545	1 SYFB_METAC	O8tpf7 methanosarc
21	39	47.0	545	1 SYFB_METNA	O8ta5 methanosarc
22	39	47.0	880	1 SVY_BACST	P11931 bacillus st
23	39	47.0	1967	1 YGSO_YEAST	P53227 saccharomyc
24	38.5	46.4	172	1 DEFL_RHIME	Q92qd0 rhizobium m
25	38.5	46.4	468	1 SELA_PSEAE	Q9hv01 pseudomonas
26	38	45.8	87	1 Y152_DREPA	O9pqz2 ureaplasma
27	38	45.8	185	1 RRE_THEMA	O9x1b9 thermotoga
28	38	45.8	198	1 B1M_HUMAN	O43521 homo sapien
29	38	45.8	251	1 KDKA_VIECH	Q9xvb9 vibrio chol
30	38	45.8	384	1 ODP2_MYCPE	P47514 mycoplasma
31	38	45.8	402	1 ODP2_MYCPN	P75392 mycoplasma
32	38	45.8	432	1 TIG_ECOL6	O8fka7 escherichia
33	38	45.8	432	1 TIG_ECOLI	P22257 escherichia

RESULT 1

ID	BAD_HUMAN	STANDARD;	PRT;	168 AA.
AC	Q92934; O14803;			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	16-OCT-2001 (Rel. 40, Last sequence update)			
DT	15-SEP-2003 (Rel. 42, Last annotation update)			
DE	Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component 6) (Bcl-XL/Bcl-2 associated death promoter) (BCL2-like 8 protein).			
GN	BAD OR BCL6 OR BCL2L8.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	Yin D.X., Li Z., Huang B., Chen S., Zhou H.;			
RT	"A human protein that interacts with Bcl-2 and have homology to mouse			
RT	BAD.";			
RL	Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.			
RN	[2]			
RP	SEQUENCE FROM N.A., AND PHOSPHORYLATION BY RAF-1.			
RX	MEDLINE=97083574; PubMed=8929532;			
RA	Wang H.-G., Rapp U.R., Reed J.C.;			
RT	"Bcl-2 targets the protein Kinase Raf-1 to mitochondria.";			
RL	Cell 87:629-638(1996).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RA	Takayama S., Reed J.C.;			
RT	Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.			
RN	[4]			
RP	SEQUENCE FROM N.A., AND DIMERIZATION.			
RC	TISSUE=Bone marrow;			
RX	MEDLINE=98049554; PubMed=9388232;			
RA	Ottlie S., Diaz J.-L., Horne W., Chang J., Wang Y., Wilson G.,			
RT	Chang S., Weeks S., Fritz L.C., Oltersdorf T.;			
RL	"Dimerization properties of human BAD.";			
RN	[5]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Lung;			
RX	MEDLINE=22388257; PubMed=12477932;			
RA	Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,			
RA	Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,			
RA	Altshul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,			
RA	Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,			
RA	Diatchenko L., Marsina K., Farmer A.A., Rubin G.M., Hong L.,			
RA	Stapleton M., Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,			
RA	Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,			
RA	Raha S.S., Lequellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,			
RA	Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,			
RA	Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,			
RA	Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,			
RA	Raheij J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,			
RA	Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,			
RA	Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,			

ALIGNMENTS

Q9n0w2 b alpha-(1,
Q9byc3 h alpha-(1,
P79282 s alpha-(1,
Q816y4 arabidopsis
P34703 caenorhabdi
Q18823 caenorhabdi
P24117 mycoplasma
P75728 escherichia
O28303 archaeoglob
Q55587 synechocyst
Q99728 homo sapien
Q49640 mycobacteri

34 38 45.8 575 1 FUT8_BOVIN
35 38 45.8 575 1 FUT8_HUMAN
36 38 45.8 575 1 FUT8_PIG
37 38 45.8 631 1 EMF2_ARATH
38 38 45.8 1521 1 EMB5_CABEL
39 38 45.8 1535 1 LML1_CABEL
40 37 44.6 375 1 DP3B_MYCCA
41 37 44.6 391 1 UBIF_ECOLI
42 37 44.6 398 1 PSMR_ARCFU
43 37 44.6 481 1 Y335_SYNY3
44 37 44.6 777 1 BAR1_HUMAN
45 37 44.6 787 1 REIA_MYCLE

RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalilus D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [6].
 RP STRUCTURE BY NMR OF 103-127.
 RX MEDLINE=21073561; PubMed=11206074;
 RA Petros A.M., Nettesheim D.G., Wang Y., Olejniczak E.T., Meadows R.P.,
 RA Mack J., Swift K., Matayoshi E.D., Zhang H., Thompson C.B.,
 RA Pesik S.W.;
 RT "Rationale for Bcl-xL/Bad peptide complex formation from structure,
 RT mutagenesis, and biophysical studies";
 RL Protein Sci. 9:2528-2534(2000).
 CC -!- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2 (By
 CC similarity). Appears to act as a link between growth factor
 CC receptor signaling and the apoptotic pathways.
 CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 CC The Ser-75/Ser-99 phosphorylated form binds 14-3-3 proteins (By
 CC similarity).
 CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm.
 CC -!- TISSUE SPECIFICITY: EXPRESSED IN A WIDE VARIETY OF TISSUES.
 CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -!- PTM: Phosphorylated on Ser-75 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-99 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-118, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-99 is the major site of AKT/PKB phosphorylation, Ser-118 the
 CC major site of protein kinase A (CAK) phosphorylation (by
 CC similarity).
 CC -!- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
 CC -!- SIMILARITY: BELONGS TO THE BCL-2 FAMILY.
 CC -!- CAUTION: Ref.1 sequence differs from that shown due to frameshifts
 CC in position 64 and 91.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: U66879; AAB36515.1; ALT_FRAME.
 DR EMBL: AF021792; AAB72092.1; -.
 DR EMBL: AF031523; AAB88124.1; -.
 DR EMBL: BC001901; AAB01901.1; -.
 DR PDB: 1G5J; 07-FEB-01.
 DR Genew: HGNC:936; BAD.
 DR MTM: 603167; -.
 DR GO: GO:0005737; Cytoplasm; NAS.
 DR GO: GO:0005741; C:mitochondrial outer membrane; NAS.
 DR GO: GO:0005515; F:protein binding activity; NAS.
 DR GO: GO:0008632; P:apoptotic program; TAS.
 DR GO: GO:0006917; P:induction of apoptosis; NAS.
 DR InterPro: IPR000712; Bcl2 BH.
 DR PROSITE: PS01259; BH3; FALSE_NEG.
 DR Apoptosis; Phosphorylation; Polymorphism; 3D-structure.
 KW DOMAIN 110 124 BH3.
 FT MOD_RES 75 75 PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).
 FT MOD_RES 99 99 PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).
 FT MOD_RES 118 118 PHOSPHORYLATION (BY PKA AND PKB) (BY
 FT SIMILARITY).

FT FT SIMILARITY).
 FT A -> S (in dbSNP:3729933).
 FT /FTId=VAR_015380.
 FT HELIX 106 121
 SQ SEQUENCE 168 AA; 18392 MW; 69FD8D27DDEE3241 CRC64;
 Query Match 100.0%; Score 83; DB 1; Length 168;
 Best Local Similarity 100.0%; Pred. No. 1.8e-06;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QRYGRLRRMSDFVD 16
 DQ |||||
 Db 108 QRYGRLRRMSDFVD 123
 RESULT 2
 ID BAD_MOUSE STANDARD; PRT; 204 AA.
 AC Q61337;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DE 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Bcl2-antagonist of cell death (BAD) (Bcl-2 binding component
 DE 6) (Bcl-xL/Bcl-2 associated death promoter).
 GN BAD OR BCL2.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RC SEQUENCE FROM N.A.
 RP TISSUE=Brain, and Thymus;
 RX MEDLINE=95136361; PubMed=7834748;
 RA Yang E., Zha J., Jockel J., Boise L.H., Thompson C.B., Korsmeyer S.J.;
 RT "Bad, a heterodimeric partner for Bcl-xL and Bcl-2, displaces Bax and
 RT promotes cell death.";
 RL Cell 80:285-291(1995).
 RN [2]
 RP PHOSPHORYLATION, AND MUTAGENESIS OF SER-112 AND SER-136.
 RX MEDLINE=98022383; PubMed=9381178;
 RA Del Peso L., Gonzalez-Garcia M., Page C., Herrera R., Nunez G.;
 RT "Interleukin-3-induced phosphorylation of BAD through the protein
 RT kinase Akt.";
 RL Science 278:687-689(1997).
 RN [3]
 RP MUTAGENESIS OF SERINE RESIDUES.
 RX MEDLINE=20403302; PubMed=10949026;
 RA Datta S.R., Katsov A., Hu L., Petros A., Pesik S.W., Yaffe M.B.,
 RA Greenberg M.E.;
 RT "14-3-3 proteins and survival kinases cooperate to inactivate BAD by
 RT BH3 domain phosphorylation.";
 RL Mol. Cell 6:41-51(2000).
 CC -!- FUNCTION: Promotes cell death. Successfully competes for the
 CC binding to Bcl-x(L), Bcl-2 and Bcl-w, thereby affecting the level
 CC of heterodimerization of these proteins with BAX. Can reverse the
 CC death repressor activity of Bcl-x(L), but not that of Bcl-2.
 CC Appears to act as a link between growth factor receptor signaling
 CC and the apoptotic pathways.
 CC -!- SUBUNIT: Forms heterodimers with the anti-apoptotic proteins, Bcl-
 CC x(L), Bcl-2 and Bcl-w. Also binds protein S100A10 (By similarity).
 CC The Ser-112/Ser-136 phosphorylated form binds 14-3-3 proteins.
 CC -!- SUBCELLULAR LOCATION: Outer mitochondrial membrane. Upon
 CC phosphorylation, locates to the cytoplasm.
 CC -!- DOMAIN: Intact BH3 domain is required by BIK, BID, BAK, BAD AND
 CC BAX for their pro-apoptotic activity and for their interaction
 CC with anti-apoptotic members of the Bcl-2 family.
 CC -!- PTM: Phosphorylated on Ser-112 in response to survival stimuli.
 CC Subsequent phosphorylation on Ser-136 promotes heterodimerization
 CC with 14-3-3 proteins. This interaction then facilitates the
 CC phosphorylation at Ser-155, a site within the BH3 domain, leading
 CC to the release of Bcl-x(L) and the promotion of cell survival.
 CC Ser-136 is the major site of AKT/PKB phosphorylation, Ser-155 the
 CC major site of protein kinase A (CAK) phosphorylation.

Best Local Similarity 100.0%; Pred. No. 9e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDEF 14
| | | | | | | | | | | | | | | | | |
Db 146 QRYGRLRMSDEF 159

RESULT 4

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BIM_MOUSE
ID BIM_MOUSE STANDARD; PRT; 196 AA.
AC 054918; 054919; 054920;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE BCL2-like protein 11 (BCL2 interacting mediator of cell death).
GN BCL2L1L OR BIM.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A., FUNCTION, SUBCELLULAR LOCATION, TISSUE
RP SPECIFICITY, AND ALTERNATIVE SPLICING.
RX MEDLINE=98094360; PubMed=9430630;
RA O'Connor L., Strasser A., O'Reilly L.A., Hausmann G., Adams J.M.,
RA Cory S., Huang D.C.S.;
RT "Bim: a novel member of the Bcl-2 family that promotes apoptosis.";
RL EMBO J. 17:384-395(1998).
CC -1- FUNCTION: INDUCES APOPTOSIS. THE ISOFORMS VARY IN CYTOTOXICITY
CC WITH ISOFORM BIM BEING THE MOST POTENT AND ISOFORM BIMEL BEING
CC THE LEAST POTENT.
CC -1- SUBUNIT: FORMS HETERODIMERS WITH A NUMBER OF ANTIAPOPTOTIC BCL-2
CC PROTEINS INCLUDING MCL-1, BCL-2, BCL-XL, BFL-1, AND BHRF-1. DOES
CC NOT HETERODIMERIZE WITH PROAPOPTOTIC PROTEINS SUCH AS BAD, BOK,
CC BAX OR BAK (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: ASSOCIATED WITH INTRACYTOSOLIC MEMBRANES.
CC Event-Alternative products:
CC Name=BimEL;
CC Name=BimL;
CC Name=BimS;
CC IsoId=054918-1; Sequence=VSP_000536;
CC IsoId=054918-2; Sequence=VSP_000537;
CC IsoId=054918-3; Sequence=VSP_000537;
CC -1- TISSUE SPECIFICITY: EXPRESSED IN A NUMBER OF B-AND T-LYMPHOID CELL
CC LINES.
CC -1- DOMAIN: THE BCL2 DOMAIN IS REQUIRED FOR BCL-2 BINDING AND
CC CYTOTOXICITY.
CC -1- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF032459; AAC40029.1; -
CC EMBL; AF032460; AAC40030.1; -
CC EMBL; AF032461; AAC40031.1; -
CC MGD; MGI:1197519; Bcl2l1l.
CC InterPro; IPR000712; Bcl2_BH
CC PROSITE; PS01259; BH3; FALSE_NEG.
KW Apoptosis; Alternative splicing; Membrane.
FT DOMAIN 146 160 BH3.
FT VARSPPLIC 42 97 Missing (in isoform BimL).
FT /FTId=VSP_000536.
FT VARSPPLIC 42 127 Missing (in isoform BimS).
FT /FTId=VSP_000537.
SQ SEQUENCE 196 AA; 22066 MW; 531C176E3F1AC9AA CRC64;

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Query Match 50.6%; Score 42; DB 1; Length 196;
Best Local Similarity 61.5%; Pred. No. 8.7;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 RYGLRMSDEF 14
| | | | | | | | | | | | | | | | | |
Db 145 RYGLRMSDEF 157

RESULT 5

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BIM_RAT
ID BIM_RAT STANDARD; PRT; 196 AA.
AC 088498; 088497; 09WU18;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE BCL2-like protein 11 (BCL2 interacting mediator of cell death)
GN BCL2L1L OR BIM OR BOD.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A., FUNCTION, SUBUNIT, AND TISSUE SPECIFICITY
RP (ISOFORMS BOD-L; BOD-M AND BOD-S).
RX MEDLINE=98400436; PubMed=9731710;
RA Hsu S.Y., Lin P.Y., Hsueh A.J.W.;
RT "BOD (Bcl-2-related ovarian death gene) is an ovarian BH3 domain-
RT containing proapoptotic Bcl-2 protein capable of dimerization with
RT diverse antiapoptotic Bcl-2 members.";
RL Mol. Endocrinol. 12:1432-1440(1998).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM BIML).
RA Chen D., Simon R.P., Chen J.;
RA "Cloning of rat bimL and bimL, and their differential expression in
RA ischemia and normal rat brain.";
RL Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INDUCES APOPTOSIS.
CC -1- SUBUNIT: FORMS HETERODIMERS WITH A NUMBER OF ANTIAPOPTOTIC BCL-2
CC PROTEINS INCLUDING MCL-1, BCL-2, BCL-XL, BFL-1, AND BHRF-1. DOES
CC NOT HETERODIMERIZE WITH PROAPOPTOTIC PROTEINS SUCH AS BAD, BOK,
CC BAX OR BAK.
CC -1- SUBCELLULAR LOCATION: ASSOCIATED WITH INTRACYTOSOLIC MEMBRANES
CC (BY SIMILARITY).
CC Event-Alternative products:
CC Name=BOD-L;
CC IsoId=088498-1; Sequence=Displayed;
CC Note=Isoform BOD-S is produced by alternative initiation at
CC Met-104 of isoform BOD-L;
CC Name=BimL;
CC IsoId=088498-2; Sequence=VSP_000538;
CC Name=BOD-M;
CC IsoId=088498-3; Sequence=VSP_000539;
CC Event-Alternative initiation:
CC Comment=2 isoforms BOD-L (shown here) and BOD-S, are produced
CC by alternative initiation at Met-1 and Met-104;
CC -1- TISSUE SPECIFICITY: Widely expressed.
CC -1- DOMAIN: THE BH3 DOMAIN IS REQUIRED FOR BCL-2 BINDING AND
CC CYTOTOXICITY.
CC -1- SIMILARITY: Contains 1 Bcl-2 homology 3 (BH3) domain.
CC
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF065433; AAC23595.1; -

```

DR EMBL; AF065431; AAC23593.1; -;
 DR EMBL; AF065432; AAC23594.1; -;
 DR EMBL; AF136927; AAD26594.1; -;
 DR InterPro; IPR000712; Bcl2_BH.
 KW Apoptosis; Alternative splicing; Membrane; Alternative initiation.
 FT CHAIN 1 196 BCL2-LIKE PROTEIN 11, ISOFORM BOD-L.
 FT CHAIN 104 196 BCL2-LIKE PROTEIN 11, ISOFORM BOD-S.
 FT INIT_MET 104 104 FOR ISOFORM BOD-S.
 FT DOMAIN 146 160 BH3.
 FT VARSPLIC 42 97 Missing (in isoform Bimt).
 FT VARSPLIC 42 127 /FTid=VSP_000538.
 FT VARSPLIC 136 136 Missing (in isoform BOD-M).
 FT CONFLICT 136 136 /FTid=VSP_000539.
 FT CONFLICT 136 136 E -> D (IN REF. 1; AAC23594).
 SQ SEQUENCE 196 AA; 22055 MW; B4D2146F9C0B37A0 CRC64;
 Query Match 50.6%; Score 42; DB 1; Length 196;
 Best Local Similarity 61.5%; Pred. No. 8.7;
 Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 2 RYGRLERRMSDF 14
 I :|||: |||
 Db 145 RIQLRLRGDF 157
 RESULT 6
 HGD_RHILO
 ID HGD_RHILO STANDARD; PRT; 457 AA.
 AC Q983J4;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Homogentisate 1,2-dioxygenase (EC 1.13.11.5) (Homogentisicase)
 DE (Homogentisate oxygenase) (Homogentisic acid oxidase).
 GN HMG A OR MLR8303.
 OS Rhizobium loti (Mesorhizobium loti).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
 OC Phyllobacteriaceae; Mesorhizobium.
 OX NCBI_TaxID=381;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MAFF303099;
 RX MEDLINE=21082930; PubMed=11214968;
 RA Kaneo T., Nakamura Y., Sato S., Asamizu E., Kato T., Sasamoto S.,
 RA Watanabe A., Iidesawa K., Ishikawa A., Kawashima K., Kimura T.,
 RA Kishida Y., Kiyokawa C., Kohara M., Matsumoto M., Matsuno A.,
 RA Mochizuki Y., Nakayama S., Nakazaki N., Shimpo S., Sugimoto M.,
 RA Takeuchi C., Yamada M., Tabata S.;
 RT "Complete genome structure of the nitrogen-fixing symbiotic bacterium
 Mesorhizobium loti." (2000).
 RL DNA Res. 7:331-338(2000).
 CC -!- CATALYTIC ACTIVITY: Homogentisate + O(2) = 4-maleylacetoacetate.
 CC -!- COFACTOR: Iron (by similarity).
 CC -!- PATHWAY: Catabolism of tyrosine; third step.
 CC -!- PATHWAY: Catabolism of phenylalanine; fourth step.
 CC -!- SIMILARITY: Belongs to the homogentisate dioxygenase family.
 CC
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 CC -----
 DR EMBL; AP030133; BAB53887.1; -;
 DR HAMAP; MF_00334; -; 1.
 DR InterPro; IPR005708; HmgA.
 DR Pfam; PF04209; HmgA; 1.
 DR TIGRFAMs; TIGR01015; hmgA; 1.
 KW Oxidoreductase; Dioxygenase; Phenylalanine catabolism;
 KW Tyrosine catabolism; Metal-binding; Iron; Complete proteome.

FT METAL 351 351 IRON (BY SIMILARITY).
 FT METAL 357 357 IRON (BY SIMILARITY).
 FT METAL 387 387 IRON (BY SIMILARITY).
 SQ SEQUENCE 457 AA; 51046 MW; 6A20B69E9A2B2BD1 CRC64;
 Query Match 50.6%; Score 42; DB 1; Length 457;
 Best Local Similarity 46.7%; Pred. No. 21;
 Matches 7; Conservative 2; Mismatches 6; Indels 0; Gaps 0;
 QY 2 RYGRLERRMSDFVD 16
 ||| || | :|
 Db 425 RYGALETRQDNYID 439
 RESULT 7
 HT2A_HUMAN
 ID HT2A_HUMAN STANDARD; PRT; 653 AA.
 AC Q13049; O9NOP8;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Zinc-finger protein HT2A (72 kDa Tat-interacting protein) (Tripartite
 DE motif-containing protein 32).
 GN TRIM32 OR HT2A.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95297135; PubMed=7778269;
 RA Fridell R.A., Harding L.S., Bogerd L.S., Boyer H.P., Cullen B.R.;
 RT "Identification of a novel human zinc finger protein that
 RT specifically interacts with the activation domain of lentiviral Tat
 RT proteins.";
 RL Virology 209:347-357(1995).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Sehra H.;
 RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Skin.
 RX MEDLINE=22386257; PubMed=12477932;
 RA Strausberg R.L., Feilgold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altshul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Ustin J.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villallon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blacksteley K.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butlerfield Y.S.N., Krzywinski M.I., Skalska U., Smalish D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 CC -!- FUNCTION: MAY PLAY A SIGNIFICANT ROLE IN MEDIATING THE BIOLOGICAL
 CC ACTIVITY OF THE HIV-1 TAT PROTEIN IN VIVO. BINDS SPECIFICALLY TO
 CC THE ACTIVATION DOMAIN OF HIV-1 TAT AND CAN ALSO INTERACT WITH THE
 CC HIV-2 AND ELAV TAT PROTEINS IN VIVO.
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- TISSUE SPECIFICITY: SPLEEN, THYMUS, PROSTATE, TESTIS, OVARY,
 CC INTESTINE AND COLON.
 CC -!- SIMILARITY: Contains 1 RING-type zinc finger.
 CC -!- SIMILARITY: Contains 1 B box-type zinc finger.

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 CC -----
 CC EMBL; U18543; AAA86474.1; -;
 CC EMBL; ALL33284; CAB92723.1; -;
 CC EMBL; BC003154; AA03154.1; -;
 CC HSP; P29590; IBOR
 CC Genew; HGNC:16380; TRM32.
 CC MIM; 602290; -;
 CC GO; GO:0005634; C:nucleus; TAS.
 CC GO; GO:0003713; F:transcription co-activator activity; TAS.
 CC InterPro; IPR001258; NHL.
 CC InterPro; IPR000315; Znf.Bbox.
 CC InterPro; IPR001841; Znf_ring.
 CC Pfam; PF01436; NHL; 5.
 CC Pfam; PF00643; zf-B_box; 1.
 CC Pfam; PF00097; zf-C3HC4; 1.
 CC SMART; SM00336; BBOX; 1.
 CC SMART; SM00184; RING; 1.
 CC PROSITE; PS00119; ZF_BBOX; 1.
 CC PROSITE; PS00518; ZF_RING_1; 1.
 CC PROSITE; PS00089; ZF_RING_2; 1.
 CC Zinc-finger; Nuclear protein.
 CC DOMAIN 2 6 POLY-ALA.
 CC ZN_RING 20 65 RING-TYPE.
 CC ZN_RING 103 133 B_BOX-TYPE.
 CC FT ZN_RING 27 27 F -> I (IN REF. 1).
 CC FT CONFLICT 27 27
 CC SEQUENCE 653 AA; 71988 MW; D83B1595CA8578FD CRC64;
 KW Aromatic amino acid biosynthesis; Lysase; Multigene family.
 SQ

Query Match 50.6%; Score 42; DB 1; Length 653;
 Best Local Similarity 61.5%; Pred. No. 30;
 Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDE 13
 Db 186 QYGEHRRYQDE 198

RESULT 8
 ID AROG_CANAL STANDARD; PRT; 370 AA.
 AC P79023;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Phospho-2-dehydro-3-deoxyheptonate aldolase, tyrosine-inhibited
 DE (EC 4.1.2.15) (Phospho-2-keto-3-deoxyheptonate aldolase) (DAMP
 DE synthetase) (3-deoxy-D-arabino-heptulosonate 7-phosphate synthase).
 GN ARO4.
 OS Candida albicans (Yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; mitosporic Saccharomycetales; Candida.
 OX NCBI_TaxID=5476;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA STRAIN=ATCC 11651 / B792;
 RA Sousa S., Pereira S.A., Livi G.P.;
 RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP PARTIAL SEQUENCE FROM N.A.
 RA MEDLINE=96207468; PubMed=8625423;
 RA Pereira S.A., Livi G.P.;
 RT "Aromatic amino-acid biosynthesis in Candida albicans: identification
 RT of the ARO4 gene encoding a second DHP synthase."
 RL Curr. Genet. 29:441-445(1996).
 CC -!- FUNCTION: STEREOSPECIFIC CONDENSATION OF PHOSPHOENOLPYRUVATE (PEP)
 CC AND D-ERYTHROSE-4-PHOSPHATE (BAP) GIVING RISE TO 3-DEOXY-D-

CC ARABINO-HEPTULOSONATE-7-PHOSPHATE (DAMP).
 CC -!- CATALYTIC ACTIVITY: 2-dehydro-3-deoxy-D-arabino-heptonate 7-
 CC phosphate + phosphate = phosphoenolpyruvate + D-erythrose 4-
 CC phosphate + H(2)O.
 CC -!- ENZYME REGULATION: INHIBITED BY TYROSINE (BY SIMILARITY).
 CC -!- PATHWAY: Aromatic amino acids biosynthesis; shikimate pathway;
 CC first step.
 CC -!- SIMILARITY: BELONGS TO CLASS-I DHP SYNTHETASE FAMILY.
 CC -----
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 CC -----
 CC EMBL; U53216; AAB48240.1; -;
 CC HSP; P00886; IQR7
 CC InterPro; IPR006219; AROFGH.
 CC Pfam; PF00793; DHP_synth_1; 1.
 CC ProDom; PD005060; AROFGH; 1.
 CC TIGRFAMs; TIGR00034; aroFGH; 1.
 CC Aromatic amino acid biosynthesis; Lysase; Multigene family.
 SQ SEQUENCE 370 AA; 40291 MW; 11E5E324C8D7B6DB CRC64;
 KW Aromatic amino acid biosynthesis; Lysase; Multigene family.
 SQ

Query Match 49.4%; Score 41; DB 1; Length 370;
 Best Local Similarity 42.9%; Pred. No. 24;
 Matches 6; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRELRMSDEFVD 16
 Db 77 YGRLKLKLADELKD 90

RESULT 9
 GCR3_YEAST STANDARD; PRT; 861 AA.
 ID GCR3_YEAST
 AC F34160;
 DT 01-FEB-1994 (Rel. 28, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE GCR3 protein (STO1 protein) (SUT1 protein).
 GN GCR3 OR STO1 OR SUT1 OR YMR125W OR YMR564.07 OR YMR553.01.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=92380925; PubMed=1512188;
 RA Oemura H., Jigami Y.;
 RT "GCR3 encodes an acidic protein that is required for expression of
 RT glycolytic genes in Saccharomyces cerevisiae."
 RL J. Bacteriol. 174:5526-5532(1992).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Pandit S., Sternglanz R.;
 RA Submitted (DEC-1992) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Guo Z., Russo P., Sherman F.;
 RA Submitted (XXX-1994) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RA STRAIN=S288c / AB972;
 RA PubMed=9169872;
 RA Bowman S., Churcher C.M., Badcock K., Brown D., Chillingworth T.,
 RA Connor R., Dedman K., Devlin K., Gentles S., Hamlin N., Hunt S.,
 RA Jagels K., Lye G., Moule S., Odell C., Pearson D., Rajandream M.A.,
 RA Rice P., Skelton J., Walsh S., Whitehead S., Barrell B.G.;
 RT "The nucleotide sequence of Saccharomyces cerevisiae chromosome

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RT XIII."
RL Nature 387:90-93(1997).
CC -!- FUNCTION: REQUIRED FOR EXPRESSION OF GLYCOLYTIC GENES. HAS
CC CERTAIN CHARACTERISTICS OF A TRANSCRIPTIONAL ACTIVATOR.
CC -!- SUBCELLULAR LOCATION: Nuclear (Probable).
CC -!- SIMILARITY: SOME, TO HUMAN CBP80
CC -!- SIMILARITY: REF.3 SEQUENCE DIFFERS FROM THAT SHOWN FROM POSITION 708
CC ONWARD AND IS SHORTER (725 AA) DUE TO A FRAMESHIFT.
CC -----
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CC -----
DR EMBL: D10224; BAA01076.1; ALT SEQ.
DR EMBL: L07650; -; NOT_ANNOTATED_CDS.
DR EMBL: L27744; -; NOT_ANNOTATED_CDS.
DR EMBL: Z49273; CAA89274.1; -.
DR EMBL: Z48622; CAA8850.1; -.
DR PIR: A44919; A44919.
DR SGD: S0004732; STOL.
DR GO: GO:0000243; C:commitment complex; IPI.
DR GO: GO:0005846; C:sRNA cap binding complex; IDA.
DR GO: GO:0003729; F:mRNA binding activity; IPI.
DR GO: GO:0006371; P:mRNA splicing; IPI.
DR InterPro: IPR003890; IF_eIF4G.
DR Pfam: PF02854; MIF4G; 1.
DR SMART: SM00543; MIF4G; 1.
KW DNA-binding; Nuclear protein.
FT DOMAIN 22 30 NUCLEAR LOCALIZATION SIGNAL (POTENTIAL).
FT DOMAIN 774 801 ASP/GLU-RICH (ACIDIC).
FT DOMAIN 802 825 ARG/LYS-RICH (BASIC).
FT CONFLICT 164 164 D -> V (IN REF. 3).
FT CONFLICT 633 633 R -> I (IN REF. 3).
FT CONFLICT 704 704 A -> R (IN REF. 3).
FT SEQUENCE 861 AA; 100017 MW; EDD04907BDC9207D CRC64;
Query Match 49.4%; Score 41; DB 1; Length 861;
Best Local Similarity 40.0%; Pred. No. 58;
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 1 QRYGRLRMSDEFV 15
Db 821 RRYSHYRELADKFI 835

RESULT 10
YGFGL_ECOLI
ID YGFGL_ECOLI STANDARD; PRT; 261 AA.
AC P52045; P76643;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hypothetical protein ygfG.
GN YGFGL OR B2919.
OS Escherichia coli.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blatter F.R., Plunkett G., III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12."
RL Science 277:1453-1474(1997).
CC -!- SIMILARITY: BELONGS TO THE ENOYL-COA HYDRATASE/ISOMERASE FAMILY.

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CC -----
DR EMBL: U28377; AAA69086.1; ALT INIT.
DR EMBL: AAC00375; AAC75956.1; ALT_INIT.
DR PDB: 1ER8; 24-MAY-00.
DR PDB: 1EP9; 24-MAY-00.
DR Ecogene; EGI2972; YgfG.
DR InterPro: IPR001753; EnCoA_hydrase.
DR Pfam: PF00378; ECH; 1.
DR PROSITE; PS00166; ENOYL_COA_HYDRATASE; 1.
KW Hypothetical protein; Lyase; Complete proteome; 3D-structure.
SQ SEQUENCE 261 AA; 29172 MW; B6A8A13EC2C2EBE0 CRC64;
Query Match 48.2%; Score 40; DB 1; Length 261;
Best Local Similarity 50.0%; Pred. No. 25;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 3 YGRLRMSDEFVD 16
Db 21 YGRKLNLSKVFIID 34

RESULT 11
PHLC_TRYCR
ID PHLC_TRYCR STANDARD; PRT; 380 AA.
AC O15886;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Variant-surface-glycoprotein phospholipase C (EC 3.1.4.47) (VSG
DE lipase) (Glycosylphosphatidylinositol-specific phospholipase C)
DE (GPI-PLC).
OS Trypanosoma cruzi.
OC Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.
OX NCBI_TaxID=5693;
RN [1]
RP SEQUENCE FROM N.A.
RA Redpath M., Carnall N., Webb H., Courel M., Amorim A.,
RA Cardosoalmeida M.L., Carrington M.;
RA Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: BY HYDROLYSIS OF THE ATTACHED GLYCOLIPID, RELEASES
CC SOLUBLE VARIANT SURFACE GLYCOPROTEIN CONTAINING PHOSPHOINOSITOL
CC FROM THE CELL WALL OF T.BRUEI AFTER CELL LYSIS. IT ALSO CLEAVES
CC SIMILAR MEMBRANE ANCHORS ON SOME MAMMALIAN PROTEINS. VSG LIPASE
CC MAY PLAY A ROLE IN PROCESSES SUCH AS PARASITE DIFFERENTIATION OR
CC ANTIGENIC VARIATION (BY SIMILARITY).
CC -!- CATALYTIC ACTIVITY: Variant-surface-glycoprotein 1,2-didecanoyl-
CC sn-phosphatidylinositol + H(2)O = 1,2-didecanoylglycerol + soluble
CC variant-surface-glycoprotein.
CC -!- SUBUNIT: Monomer (By similarity).
CC -!- SUBCELLULAR LOCATION: Membrane-associated.
CC -!- SIMILARITY: DOMAIN X IS CONSERVED IN DIFFERENT FORMS OF PLC AND IS
CC ESSENTIAL FOR CATALYTIC ACTIVITY.
CC -----
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CC -----
DR EMBL: AJ000079; CAA03904.1; -.
DR InterPro: IPR000909; PI-PLC_xdom.
DR InterPro: IPR003633; Varsurfglyc_PLIC.
DR Pfam: PF00388; PI-PLC-X; 1.

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DR Pfam: PF03490; Varsurf_PPLC; 1.
 DR ProDom: PD041675; Varsurfglyc_PPLC; 1.
 DR SMART: SM00148; PLCX; 1.
 DR PROSITE: PS00007; PPLC_X_DOMAIN; 1.
 KW HydroLase; Membrane.
 FT DOMAIN 31 205 DOMAIN X.
 SQ SEQUENCE 380 AA; 42736 MW; 273CD402B52068C5 CRC64;

Query Match 48.2%; Score 40; DB 1; Length 380;
 Best Local Similarity 50.0%; Pred. No. 36;
 Matches 7; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 2 RYGRLEIRMSDEFV 15
 : ||| I: || I:
 DB 165 KFFELDLRLSDRF 178

RESULT 12
 MTAL ACBPA STANDARD; PRT; 429 AA.
 AC O52702;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Modification methylase Apali (EC 2.1.1.73) (Cytosine-specific
 methyltransferase Apali) (M.Apali).
 GN APALI.
 OS Acetobacter pasteurianus (Acetobacter turbidans).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodospirillales;
 OC Acetobacteraceae; Acetobacter.
 OX NCBI_taxid=438;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-ATCC 12875;
 RX MEDLINE-99077292; PubMed=9862476;
 RA Xu S.-Y., Xiao J.-P., Ettwiller L., Holden M., Aliotta J., Poh C.I.,
 Dalton M., Robinson D.P., Petronzio T.R., Moran L., Ganatra M.,
 Ware J., Slatko B., Benner J. II;
 RT Cloning and expression of the Apali, NspI, NspH, SaeI, SaeI, and
 RT Sapi restriction-modification systems in *Escherichia coli*.;
 RL Mol. Gen. Genet. 260:228-231(1998).
 CC -!- FUNCTION: THIS METHYLASE RECOGNIZES THE DOUBLE-STRANDED SEQUENCE
 CC GTCCAC, CAUSES SPECIFIC METHYLATION ON C-? ON BOTH STRANDS, AND
 CC PROTECTS THE DNA FROM CLEAVAGE BY THE APALI ENDONUCLEASE.
 CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + DNA cytosine = S-
 CC adenosyl-L-homocysteine + DNA 5-methylcytosine.
 CC -!- SIMILARITY: BELONGS TO THE C5-METHYLTRANSFERASE FAMILY.

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 CC -----
 DR EMBL: AF044847; AAC97180.1;
 DR REBASE: 3281; M.APALI
 DR InterPro: IPR001525; C5_DNA_meth.
 DR Pfam: PF00145; DNA_methylase; 1.
 DR PRINTS: PRO0105; C5METHTRFASE.
 DR TIGRFAMs: TIGR00675; dcm; 1.
 DR PROSITE: PS00094; C5_MTASE1; FALSE NEG.
 DR Transferase; Methyltransferase; Restriction system.
 FT ACT_SITE 81 81 BY SIMILARITY.
 SQ SEQUENCE 429 AA; 46547 MW; E011C7D15B33F5F3 CRC64;

Query Match 48.2%; Score 40; DB 1; Length 429;
 Best Local Similarity 61.5%; Pred. No. 41;
 Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 4 GRELRMSDEFVD 16
 ||| I: ||| I:

DB 129 GRDLARLVREFVD 141

RESULT 13

RPSD_BORBU STANDARD; PRT; 631 AA.
 AC P52323;

DT 01-OCT-1996 (Rel. 34, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE RNA polymerase sigma factor rpoD (Sigma-70).
 GN RPOD OR BB0712.

OS *Borrelia burgdorferi* (Lyme disease spirochete).
 OC Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; *Borrelia*.
 OX NCBI_taxid=139;

RN [1]
 RP SEQUENCE OF 89-631 FROM N.A.
 RC STRAIN-ATCC 35210 / B31;

RL Thesis (1994), National Taiwan University, Taiwan.
 RN [2]
 RP SEQUENCE FROM N.A.

RC STRAIN-ATCC 35210 / B31;
 RX MEDLINE-98065943; PubMed=9403685;

RA Fraser C.W., Casjens S., Huang W.A., Dodson R., Hickey E.K., Gwinn M.,
 RA Lathigra R., White O., Ketchum K.A., Fleischmann R.D., Richardson D.,
 RA Dougherty B., Tomb J.-F., Fleischmann R.D., Richardson D.,
 RA Peterson J., Kerlavage A.R., Quackenbush J., Salzberg S., Hanson M.,
 RA van Vugt R., Palmer N., Adams M.D., Gocayne J.D., Weidman J.,
 RA Utterback T., Wathley L., McDonald L., Artiach P., Bowman C.,
 RA Garland S., Fujii C., Cotton M.D., Horst K., Roberts K., Hatch B.,
 RA Smith H.O., Venter J.C.;
 RT "Genomic sequence of a Lyme disease spirochete, *Borrelia*
 RT *burgdorferi*.";
 RL Nature 390:580-586(1997).
 RN [3]
 RP SEQUENCE OF 165-614 FROM N.A.

RC STRAIN=297;
 RA Pan M., Yeh J., Tsai C.;

RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: THE SIGMA FACTOR IS AN INITIATION FACTOR THAT PROMOTES
 CC ATTACHMENT OF THE RNA POLYMERASE TO SPECIFIC INITIATION SITES AND
 CC THEN IS RELEASED. THIS IS THE PRIMARY SIGMA-FACTOR OF THIS
 CC BACTERIA.

 CC -!- SIMILARITY: Belongs to the sigma-70 factor family.
 CC -----
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 CC -----

DR EMBL: U17591; AAC44104.1;
 DR EMBL: AE001171; AAC67061.1;
 DR EMBL: U68006; AAC45100.1;
 DR PIR: G70188; G70188.
 DR HSP: P00579; ISIG.

DR TIGR: BB0712;
 DR InterPro: IPR000943; Sigma_70.
 DR Pfam: PF03979; sigma70_r1.1; 1.
 DR Pfam: PF00140; sigma70_r1.2; 1.
 DR Pfam: PF04542; sigma70_r2; 1.
 DR Pfam: PF04539; sigma70_r3; 1.
 DR Pfam: PF04545; sigma70_r4; 1.
 DR PROSITE: PS00715; SIGMA70_1; 1.
 DR PROSITE: PS00716; SIGMA70_2; 1.

KW Transcription regulation; Sigma factor; DNA-directed RNA polymerase;
 FT DOMAIN 419 432 POLYMERASE CORE BINDING (POTENTIAL).
 FT DNA_BIND 589 608 H-T-H MOTIF (BY SIMILARITY).
 SQ SEQUENCE 631 AA; 73642 MW; BD565AB7D8F44796 CRC64;

```

Query Match          48.2%; Score 40; DB 1; Length 631;
Best Local Similarity 50.0%; Pred. No. 61;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 3 YGRELIRMSDEFVD 16
   I I I I I I I I I I
Db 279 YQELIRFSDDYID 292

RESULT 14
6PGL_THEME
ID 6PGL_THEME STANDARD; PRT: 220 AA.
AC O9XON8;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 6-phosphogluconolactonase (EC 3.1.1.31) (6PGL).
GN 6PGL OR DEVB OR TM1154.
OS Thermotoga maritima.
OC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
OX NCBI_TaxID=2336;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MSB8 / DSM 3109;
RX MEDLINE=99287316; PubMed=10360571;
RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
RA McDonald L., Utterback T.R., Malek J.A., Linner K.D., Garrett M.M.,
RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
RT "Evidence for lateral gene transfer between Archaea and Bacteria from
RT genome sequence of Thermotoga maritima.";
RL Nature 399:323-329(1999).
CC -!- FUNCTION: HYDROLYSIS OF 6-PHOSPHOGLUCONOLACTONE TO 6-
CC PHOSPHOGLUCONATE.
CC -!- CATALYTIC ACTIVITY: 6-phospho-D-glucono-1,5-lactone + H(2)O = 6-
CC phospho-D-gluconate.
CC -!- PATHWAY: Pentose phosphate pathway; second step.
CC -!- SIMILARITY: BELONGS TO THE GLUCOSAMINE/GALACTOSAMINE-6-PHOSPHATE
CC ISOMERASE FAMILY. 6-PHOSPHOGLUCONOLACTONASE SUBFAMILY.
CC -----
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CC -----
DR EMBL: AF001772; AAD36230.1; -
DR FICR: F72289; F72289.
DR TIGR: TM1154; -
DR InterPro: IPR006148; Gluc_gal_isom.
DR InterPro: IPR005900; Phosphoglucnolac.
DR Pfam: PF01182; Glucosamine_iso; 1.
DR TIGRfams: TIGR01198; pgl; 1.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 220 AA; 25325 MW; 9B0FD07E0C1560C3 CRC64;

Query Match          47.0%; Score 39; DB 1; Length 220;
Best Local Similarity 42.9%; Pred. No. 30;
Matches 6; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 1 QRYGELIRMSDEF 14
   : I I I I I I I I
Db 113 KYEELIRSATDQF 126

RESULT 15
RA51_SCHPO
ID RA51_SCHPO STANDARD; PRT: 365 AA.

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AC P36601;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE DNA repair protein rhp51 (RAD51 homolog).
GN RHP51 OR RAD51 OR SPAC644.14C.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=94051565; PubMed=8233794;
RA Maris D.F.R., Vreeken K., Carr A.M., Broughton B.C., Lehmann A.R.,
RA Lohman P.H.M., Pastink A.;
RT "Cloning the RAD51 homologue of Schizosaccharomyces pombe.";
RL Nucleic Acids Res. 21:4586-4591(1993).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=93364417; PubMed=8358431;
RA Shinohara A., Ogawa H., Matsuda Y., Ushio N., Ikeo K., Ogawa T.;
RT "Cloning of human, mouse and fission yeast recombination genes
RT homologous to RAD51 and recA.";
RL Nat. Genet. 4:239-243(1993).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=94255568; PubMed=8194753;
RA Jang Y.K., Jin Y.H., Kim E.M., Hong S.H., Fabre F., Park S.D.;
RT "Cloning and sequence analysis of rhp51+, a Schizosaccharomyces pombe
RT homol of the Saccharomyces cerevisiae RAD51 gene.";
RL Gene 142:207-211(1994).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21848401; PubMed=11859360;
RA Wood V., Gilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Scouras J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Feitwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Horsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skellon J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volckaert G., Aert R., Robben J., Grymonprez B.,
RA Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Moestl D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Leirach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Fumelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Domiguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Sipakovski G.V., Ussery D., Barrell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe.";
RL Nature 415:871-880(2002).
CC -!- FUNCTION: REQUIRED BOTH FOR RECOMBINATION AND FOR THE REPAIR OF
CC DNA DAMAGE CAUSED BY X-RAYS.
CC -!- SIMILARITY: BELONGS TO THE RECA FAMILY. RAD51 SUBFAMILY.
CC -----
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CC -----

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CC  EMBL; 222691; CAA80399.1; -.
DR  EMBL; D13805; BAA02963.1; -.
DR  EMBL; 224756; CAA80879.1; -.
DR  EMBL; 224756; CAA80878.1; ALT_INIT.
DR  EMBL; AL355012; CAB90141.1; -.
DR  PIR; S42107; S42107.
DR  HSSP; Q06609; 1B22.
DR  GeneDB_SPombe; SPAC64.14c; -.
DR  InterPro; IPR003593; AAA_ATPase.
DR  InterPro; IPR000445; HhH.
DR  InterPro; IPR003583; HHH_1.
DR  InterPro; IPR001553; RecA.
DR  Pfam; PF0633; HHH; 1.
DR  ProDom; PD000229; RecA; 1.
DR  SMART; SM00382; AAA; 1.
DR  SMART; SM00278; HhH1; 1.
DR  PROSITE; PS50162; RECA_2; 1.
DR  PROSITE; PS50163; RECA_3; 1.
KW  DNA damage; DNA repair; ATP-binding; DNA recombination.
FT  NP_BIND 149 156 ATP (POTENTIAL).
FT  CONFLICT 15 15 T -> M (IN REF. 2 AND 4).
SQ  SEQUENCE 365 AA; 39823 MW; 9F26E9FA4F3C2BA CRC64;

Query Match 47.0%; Score 39; DB 1; Length 365;
Best Local Similarity 53.8%; Pred No. 50;
Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RYGRRLRMSDEF 14
Db | | | | |
269 RFRMTLQRLADEF 281
```

Search completed: September 15, 2003, 17:23:01
Job time : 4.77143 secs

GenCore version 5.1.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:17:31 : Search time 17.3714 Seconds
(without alignments)
237.680 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRELRRMSDEFVD 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phase:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	59	71.1	146	13 Q919N2	Q919N2 brachydanio
2	46	55.4	458	5 Q8SV57	Q8SV57 encephalito
3	46	55.4	477	5 Q8SVF8	Q8SVF8 encephalito
4	46	55.4	641	5 Q8SU90	Q8SU90 encephalito
5	45	54.2	230	16 Q8XXS6	Q8XXS6 ralstonia s
6	45	54.2	564	16 Q9RUK9	Q9RUK9 deinococcus
7	44	53.0	247	2 Q9AJL5	Q9AJL5 streptomyce
8	44	53.0	726	16 Q8G3Z1	Q8G3Z1 bifidobacte
9	44	53.0	1248	16 Q9HZQ3	Q9HZQ3 pseudomonas
10	43	51.8	205	10 Q8L757	Q8L757 arabidopsis
11	43	51.8	804	16 Q8EW78	Q8EW78 mycoplasma
12	43	51.8	5635	5 Q9N9N1	Q9N9N1 leishmania
13	42	50.6	213	17 Q8TJ31	Q8TJ31 methanosarc
14	42	50.6	260	16 Q8RCC4	Q8RCC4 thermoaer
15	42	50.6	297	5 Q8M290	Q8M290 drosophila
16	42	50.6	339	10 Q9FQ05	Q9FQ05 atrichum an

17	42	50.6	548	2 P71029	P71029 burkholderi
18	42	50.6	592	5 Q9V716	Q9V716 drosophila
19	42	50.6	592	5 Q960B3	Q960B3 drosophila
20	42	50.6	876	16 Q931Q1	Q931Q1 staphylococ
21	42	50.6	876	16 Q99TJ8	Q99TJ8 staphylococ
22	42	50.6	876	16 Q8CS83	Q8CS83 staphylococ
23	42	50.6	990	17 Q8TUT0	Q8TUT0 methanosyru
24	41.5	50.0	191	16 Q8TG63	Q8TG63 brucella me
25	41.5	50.0	191	16 Q8GIQ8	Q8GIQ8 brucella su
26	41	49.4	216	16 Q9XZC5	Q9XZC5 streptomyce
27	41	49.4	447	10 Q9M0B1	Q9M0B1 arabidopsis
28	41	49.4	497	10 Q8L517	Q8L517 arabidopsis
29	41	49.4	577	3 Q59699	Q59699 schizosach
30	41	49.4	1303	12 Q9QB00	Q9QB00 potato mop-
31	41	49.4	2381	5 Q8IKD7	Q8IKD7 plasmodium
32	40.5	48.8	792	4 P78313	P78313 homo sapien
33	40.5	48.8	1224	4 P78311	P78311 homo sapien
34	40.5	48.8	1265	4 P78312	P78312 homo sapien
35	40	48.2	84	17 Q9HMU7	Q9HMU7 halobacteri
36	40	48.2	109	16 Q9TS19	Q9TS19 streptococc
37	40	48.2	113	16 Q8DR57	Q8DR57 streptococc
38	40	48.2	164	17 Q8ZT11	Q8ZT11 pyrobaculum
39	40	48.2	219	17 Q9UZQ0	Q9UZQ0 pyrococcus
40	40	48.2	259	17 Q8PUZ1	Q8PUZ1 methanosarc
41	40	48.2	275	16 Q8XD14	Q8XD14 escherichia
42	40	48.2	335	10 Q64692	Q64692 arabidopsis
43	40	48.2	360	16 Q910H8	Q910H8 pseudomonas
44	40	48.2	361	10 Q9FIC1	Q9FIC1 arabidopsis
45	40	48.2	411	16 Q9A3L1	Q9A3L1 caulobacter

ALIGNMENTS

RESULT 1

Q919N2 PRELIMINARY; PRT; 146 AA.
AC Q919N2;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Bad.
GN BAD.
OS Brachydanio rerio (zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20373792; PubMed=10917738;
RA Inohara N.; Nunez G.;
RT "Genes with homology to mammalian apoptosis regulators identified in zebrafish."
RL Cell Death Differ. 7:509-510(2000).
DR EMBL; AF231017; AAF66962.2; -
DR HSSP; Q29334; 1G5J
DR ZFIN; ZDB-GENE-000616-1; bad.
SQ SEQUENCE 146 AA; 16546 MW; 28A5650BB5107ECB CRC64;

Query Match 71.1%; Score 59; DB 13; Length 146;
Best Local Similarity 71.4%; Pred. No. 0.047;
Matches 10; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEF 14
:::|||||
Db 93 KRYGQGLRRMSDEF 106

RESULT 2

Q8SV57 PRELIMINARY; PRT; 458 AA.
ID Q8SV57
AC Q8SV57;

DT 01-JUN-2002 (TrEMBLrel. 21, Created)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
 DE Hypothetical protein EC006_1680.
 GN EC006_1680.
 OS Encephalitozoon cuniculi.
 OC Eukaryota; Fungi; Microsporidia; Unikaryonidae; Encephalitozoon.
 OX NCBI_TaxID=6035;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RA Genoscope;
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RX MEDLINE=21576510; PubMed=11719806;
 RA Katinka M.D., Duprat S., Cornillot E., Metenier G., Thomarat F.,
 RA Prensier G., Barbe V., Peyretailade E., Brottier P., Wincker P.,
 RA Delbac F., El Alaoui H., Peyret P., Saurin W., Gouy M.,
 RA Weissenbach J., Vivares C.P.;
 RT "Genome sequence and gene compaction of the eukaryote parasite
 Encephalitozoon cuniculi.";
 RL Nature 414:450-453(2001).
 DR EMBL; AL590446; CAD25529.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 458 AA; 51914 MW; 24111BF78DA4534D CRC64;

Query Match 55.4%; Score 46; DB 5; Length 458;
 Best Local Similarity 46.7%; Pred. No. 24;
 Matches 7; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 2 RYGRRLRMSDEFVD 16
 ||||: || | : :
 Db 13 RYGRVWRMLDDMIE 27

RESULT 3
 Q8SVF8 PRELIMINARY; PRT; 477 AA.
 AC Q8SVF8;
 DT 01-JUN-2002 (TrEMBLrel. 21, Created)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
 DE Hypothetical protein EC006_0040.
 GN EC006_0040.
 OS Encephalitozoon cuniculi.
 OC Eukaryota; Fungi; Microsporidia; Unikaryonidae; Encephalitozoon.
 OX NCBI_TaxID=6035;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RA Genoscope;
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RX MEDLINE=21576510; PubMed=11719806;
 RA Katinka M.D., Duprat S., Cornillot E., Metenier G., Thomarat F.,
 RA Prensier G., Barbe V., Peyretailade E., Brottier P., Wincker P.,
 RA Delbac F., El Alaoui H., Peyret P., Saurin W., Gouy M.,
 RA Weissenbach J., Vivares C.P.;
 RT "Genome sequence and gene compaction of the eukaryote parasite
 Encephalitozoon cuniculi.";
 RL Nature 414:450-453(2001).
 DR EMBL; AL590446; CAD25529.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 477 AA; 54039 MW; B5DCF6299724CC96 CRC64;

Query Match 55.4%; Score 46; DB 5; Length 477;
 Best Local Similarity 46.7%; Pred. No. 25;
 Matches 7; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 2 RYGRRLRMSDEFVD 16
 ||||: || | : :
 Db 32 RYGRVWRMLDDMIE 46

RESULT 4
 Q8SU90 PRELIMINARY; PRT; 641 AA.
 AC Q8SU90;
 DT 01-JUN-2002 (TrEMBLrel. 21, Created)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)
 DE Hypothetical protein EC011_0030.
 GN EC011_0030.
 OS Encephalitozoon cuniculi.
 OC Eukaryota; Fungi; Microsporidia; Unikaryonidae; Encephalitozoon.
 OX NCBI_TaxID=6035;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RA Genoscope;
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GB-MJ;
 RX MEDLINE=21576510; PubMed=11719806;
 RA Katinka M.D., Duprat S., Cornillot E., Metenier G., Thomarat F.,
 RA Prensier G., Barbe V., Peyretailade E., Brottier P., Wincker P.,
 RA Delbac F., El Alaoui H., Peyret P., Saurin W., Gouy M.,
 RA Weissenbach J., Vivares C.P.;
 RT "Genome sequence and gene compaction of the eukaryote parasite
 Encephalitozoon cuniculi.";
 RL Nature 414:450-453(2001).
 DR EMBL; AL590450; CAD25913.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 641 AA; 72408 MW; A87E6EBE4973FC2 CRC64;

Query Match 55.4%; Score 46; DB 5; Length 641;
 Best Local Similarity 46.7%; Pred. No. 34;
 Matches 7; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 2 RYGRRLRMSDEFVD 16
 ||||: || | : :
 Db 203 RYGRVWRMLDDMIE 217

RESULT 5
 Q8XXS6 PRELIMINARY; PRT; 230 AA.
 AC Q8XXS6;
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Probable ATP-binding ABC transporter protein.
 GN RSC2037 OR RS03602.
 OS Ralstonia solanacearum (Pseudomonas solanacearum).
 OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
 OC Ralstoniaceae; Ralstonia.
 OX NCBI_TaxID=305;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GMI1000;
 RX MEDLINE=21681879; PubMed=11823852;
 RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
 RA Ariat M., Billault A., Brottier P., Camus J.C., Cattolico L.,
 RA Chandler M., Choise N., Claudel-Renard C., Cunac S., Demange N.,
 RA Gaspin C., Lavie M., Moisan A., Robert C., Saurin W., Schiex T.,
 RA Siguer P., Thebault P., Whalen M., Wincker P., Levy M.,
 RA Weissenbach J., Boucher C.A.;
 RT "Genome sequence of the plant pathogen Ralstonia solanacearum.";
 RL Nature 415:497-502(2002).
 DR EMBL; AL646067; CAD15739.1; -.
 DR InterPro; IPR003593; AAA_ATPase.

DR InterPro; IPR003439; ABC_transporter.
 DR Pfam; PF00005; ABC_tran; 1.
 DR ProDom; PD000006; ABC_transporter; 1.
 DR SMART; SM00382; AAA; 1.
 KW Complete proteome.
 SQ SEQUENCE 230 AA; 25231 MW; 7C3FDALE7A19A2A4 CRC64;

Query Match 54.28; Score 45; DB 16; Length 230;
 Best Local Similarity 64.38; Pred. No. 17;
 Matches 9; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 QYGRRLRMSDEF 14
 | | | | | | | | | |
 Db 168 QEIGRTLRLRVDEF 181

RESULT 6

Q9RUK9 PRELIMINARY; PRT; 564 AA.
 AC Q9RUK9
 DT 01-MAY-2000 (TRENBLrel. 13, Created)
 DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Glycosyl hydrolase, family 13.
 GN DR1375.
 OS Deinococcus radiodurans.
 OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
 OC Deinococcaceae; Deinococcus.
 OK NCBI_TaxID=1299;
 FN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=R1;
 RX MEDLINE=20036896; PubMed=10567266;
 RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
 RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
 RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
 RA Vanthekum J.J., Lam P., McDonald L., Utterback T., Zalewski C.,
 RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
 RA Ketchum K.A., Nelson K.E., Salzberg S., Smith H.O., Venter J.C.,
 RA Fraser C.M.;
 RA "Genome sequence of the radioresistant bacterium Deinococcus
 radiodurans R1.";
 RL Science 286:1571-1577(1999).
 DR ENBL; AE001983; AAF10944.1; .
 DR HSSP; P21332; 1UOK.
 DR TIGR; DR1375; .

DR InterPro; IPR006047; Alpha_amyl_cat.
 DR InterPro; IPR006589; Alp_amyl_cat_sub.
 DR Pfam; PF00128; alpha_amylase; 1.
 DR SMART; SM00642; Aamy; 1.
 KW Hydrolase; Complete proteome.
 SQ SEQUENCE 564 AA; 63667 MW; B8F50B9B0DFC8D51 CRC64;

Query Match 54.28; Score 45; DB 16; Length 564;
 Best Local Similarity 64.38; Pred. No. 44;
 Matches 9; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRRLRMSDEFVD 16
 | | | | | | | | | |
 Db 283 YVEMRRYVDFDD 296

RESULT 7

Q9AJL5 PRELIMINARY; PRT; 247 AA.
 AC Q9AJL5
 DT 01-JUN-2001 (TRENBLrel. 17, Created)
 DT 01-JUN-2001 (TRENBLrel. 17, Last sequence update)
 DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
 DE VarR.
 GN VARR.
 OS Streptomyces virginiae.
 OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;

OC Streptomycineae; Streptomycetaceae; Streptomycetes.
 OX NCBI_TaxID=1961;
 RN [1]
 RP SEQUENCE FROM N.A.

RX MEDLINE=21125565; PubMed=11222601;
 RA Nawat W., Lee C.K., Kinoshita H., Yamada Y., Nihira T.;
 RT "Identification of the varR gene as a transcriptional regulator of
 RT virginiamycin S resistance in Streptomyces virginiae.";
 RL J. Bacteriol. 183:2025-2031(2001).
 CC -1- SIMILARITY: BELONGS TO THE TETR/ACRR FAMILY OF TRANSCRIPTIONAL
 CC REGULATORS.

DR ENBL; AB045994; BAB32408.1; .
 DR HSSP; P09164; 2TCT.
 DR InterPro; IPR001647; HTH_Tetr.
 DR InterPro; IPR004111; Tetr_C.
 DR Pfam; PF00440; tetr; 1.
 DR Pfam; PF02909; tetr_C; 1.
 DR PRINTS; PR00455; HTH_Tetr.
 DR DNA-binding; Transcription; Transcription regulation.
 KW SEQUENCE 247 AA; 27328 MW; B6326F4374598A8 CRC64;

Query Match 53.08; Score 44; DB 2; Length 247;
 Best Local Similarity 50.08; Pred. No. 27;
 Matches 7; Conservative 5; Mismatches 2; Indels 0; Gaps 0;

QY 3 YGRRLRMSDEFVD 16
 | | | | | | | | | |
 Db 186 YGKEVGRTADEFLE 199

RESULT 8

Q8G3Z1 PRELIMINARY; PRT; 726 AA.
 ID Q8G3Z1
 AC Q8G3Z1
 DT 01-MAR-2003 (TRENBLrel. 23, Created)
 DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Narrowly conserved hypothetical protein.
 GN BL1609.
 OS Bifidobacterium longum.
 OC Bacteria; Actinobacteria; Actinobacteridae; Bifidobacteriales;
 OC Bifidobacteriaceae; Bifidobacterium.
 OK NCBI_TaxID=216816;
 FN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NCC 2705;
 RX MEDLINE=22294977; PubMed=12381787;
 RA Schell M.A., Karmirantzou M., Snel B., Vilanova D., Berger B.,
 RA Pessi G., Zwahlen M.-C., Desiere F., Bork P., Delley M.,
 RA Pidmore R.D., Arigoni F.;
 RA "The genome sequence of Bifidobacterium longum reflects its adaptation
 RT to the human gastrointestinal tract.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:14422-14427(2002).
 DR ENBL; AE014794; AAN25398.1; .
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 726 AA; 81866 MW; 6BE86B61FCBC586 CRC64;

Query Match 53.08; Score 44; DB 16; Length 726;
 Best Local Similarity 60.08; Pred. No. 85;
 Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 2 YGRRLRMSDEFVD 16
 | | | | | | | | | |
 Db 241 RVGVYRRLSDEFLD 255

RESULT 9

Q9HZQ3 PRELIMINARY; PRT; 1248 AA.
 ID Q9HZQ3
 AC Q9HZQ3
 DT 01-MAR-2001 (TRENBLrel. 16, Created)
 DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Cobalamin biosynthetic protein CobN.
 GN COBN OR PA2944.

OS Pseudomonas aeruginosa.
 OC Bacteria: Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 OC Pseudomonadaceae; Pseudomonas.
 OX NCBI_TaxID=287;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=ATCC 15692 / PA01;

RX MEDLINE=20437337; PubMed=10984043;

RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,

RA Hickey M.J., Brickman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,

RA Garber R.L., Tolentino E., Westbrook-Wadman S., Yuan Y.,

RA Brady L.B., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,

RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,

RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.;

RT "Complete genome sequence of *Pseudomonas aeruginosa* PA01, an

RT opportunistic pathogen";

RL Nature 406:950-964(2000).;

DR EMBL; AB004720; A0606332.1; -

DR InterPro; IPR003672; COBN/Mg_Chitase.

DR Pfam; PF02514; COBN-Mg_chel; 1.

DR PROSITE; PS00626; RCCL_2; 1.

KW Complete proteome.

SQ SEQUENCE 1248 AA; 138499 MW; C3D3DBFEE6736C7A CRC64;

Query Match 53.08; Score 44; DB 16; Length 1248;

Best Local Similarity 50.08; Pred. No. 1.5e+02;

Matches 9; Conservative 4; Mismatches 3; Indels 2; Gaps 1;

OY 1 QRYG--RELRRMSDEYD 16

DB 615 ESYGLRDLRLADEFYD 632

RESULT 10

Q8L757

ID Q8L757 PRELIMINARY; PRT; 205 AA.

AC Q8L757;

DT 01-OCT-2002 (TRENBLrel. 22, Created)

DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Hypothetical protein.

GN A14G02715

OS Arabidopsis thaliana (Mouse-ear cress).

OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;

OC eurosids II; Brassicales; Brassicaceae; Arabidopsi.

OX NCBI_TaxID=3702;

RN [1]

RP SEQUENCE FROM N.A.

RA Southwick A., Nguyen M., Tripp M., Palm C.J., Jones T., Wu T.,

RA Carninci P., Chen H., Cheuk R., Chan M.M., Chang C.H., Dale J.M.,

RA Deng J.M., Hayashizaki Y., Huan V.W., Lee J.M., Ishida J., Kamiya A.,

RA Kawai J., Kim C.J., Narusaka M., Quach H.L., Sakurai T., Satou M.,

RA Seki M., Shinn P., Tang C.C., Toroumi M., Wallender E.K., Wong C.,

RA Wu H.C., Yamada K., Yu G., Yuan S., Shinozaki K., Ecker J.,

RA Theologis A., Davis R.W.;

RL Submitted (JUL-2002) to the EMBL/GenBank/DBSJ databases.

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=cv. Columbia;

RA Seki M., Iida K., Satou M., Sakurai T., Akiyama K., Ishida J.,

RA Nakajima M., Enju A., Kamiya A., Narusaka M., Carninci P., Kawai J.,

RA Hayashizaki Y., Shinozaki K.;

RT "Arabidopsis thaliana full-length cDNA.";

RL Submitted (NOV-2002) to the EMBL/GenBank/DBSJ databases.

DR EMBL; AY136474; AAM97139.1; -

DR EMBL; AK118269; BAC42887.1; -

KW Hypothetical protein.

SQ SEQUENCE 205 AA; 22583 MW; 1BD8D1358ECFF81 CRC64;

Query Match 51.88; Score 43; DB 10; Length 205;

Best Local Similarity 51.58; Pred. No. 32;

Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 4 GRELRMSDEYD 16

DB 100 GSQIRRCSEFYD 112

RESULT 11

Q8EW78

ID Q8EW78 PRELIMINARY; PRT; 804 AA.

AC Q8EW78;

DT 01-MAR-2003 (TRENBLrel. 23, Created)

DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Cation-transporting p-type ATPase.

GN MPPE3250.

OS Mycoplasma penetrans.

OC Bacteria; Firmicutes; Mollicutes; Mycoplasmataceae; Mycoplasma.

OX NCBI_TaxID=28227;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=HF-2;

RA MEDLINE=22354719; PubMed=12466555;

RA Sasaki Y., Ishikawa J., Yamashita A., Oshima K., Kenri T., Furuya K.,

RA Yoshino C., Horino A., Shiba T., Sasaki T., Hattori M.;

RT "The complete genomic sequence of *Mycoplasma penetrans*, an

RT intracellular bacterial pathogen in humans.";

RL Nucleic Acids Res. 30:5293-5300(2002).

DR EMBL; AF004171; BAC44118.1; -

KW Complete proteome.

SQ SEQUENCE 804 AA; 88008 MW; 69C7LAA628FF7A3 CRC64;

Query Match 51.88; Score 43; DB 16; Length 804;

Best Local Similarity 66.78; Pred. No. 1.4e+02;

Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 4 GRELRMSDEYD 15

DB 428 GSELRRMSDEYL 439

RESULT 12

Q9N9N1

ID Q9N9N1 PRELIMINARY; PRT; 5635 AA.

AC Q9N9N1;

DT 01-OCT-2000 (TRENBLrel. 15, Created)

DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Dynein heavy chain, cytosolic.

GN L3302.02

OS Leishmania major.

OC Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae; Leishmania.

OX NCBI_TaxID=5664;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=Friedlin;

RA Hilbert H., Wedler H., Wedler E., Duesterhoeft A., Ivens A.C.,

RA Quail M., Rajandream M.A., Barrell B.G.;

RL Submitted (JUN-2000) to the EMBL/GenBank/DBSJ databases.

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=Friedlin;

RX MEDLINE=98146435; PubMed=9477341;

RA Ivens A.C., Lewis S.M., Bagherzadeh A., Zhang L., Chan H.M.,

RA Smith D.F.;

RT "A physical map of the *Leishmania* major Friedlin genome.";

RL Genome Res. 8:135-145(1998).;

DR EMBL; AL359781; CAB95305.1; -

DR InterPro; IPR003593; AAA_ATPase.

DR InterPro; IPR004273; Dynein_heavy.

DR InterPro; IPR000169; Shprot_acsite.

```

DR Pfam: PF03028: Dyncin_heavy; 1.
DR SMART: SM00382; AAA; 4.
DR PROSITE: PS00639; THIOL_PROTEASE_HIS; 1.
KW ATP-binding.
SQ SEQUENCE 5635 AA; 620050 MW; 64A9E881A9B14641 CRC64;

  Query Match          51.8%; Score 43; DB 5; Length 5635;
  Best Local Similarity 72.7%; Pred. No. 1.le+03;
  Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 QRYGRELRRMS 11
   ||:|||||
Db 1535 QRYGRELRRMS 1545

RESULT 13
Q8RTJ31
ID Q8RTJ31 PRELIMINARY; PRT; 213 AA.
AC Q8RTJ31;
DT 01-JUN-2002 (TRENBLrel. 21, Created)
DT 01-JUN-2002 (TRENBLrel. 21, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE ABC transporter, ATP-binding protein.
GN NA3957.
OS Methanosarcina acetivorans.
OC Archaea; Euryarchaeota; Methanococci; Methanosarcinales;
OC Methanosarcinaceae; Methanosarcina.
OX NCBI_TaxID=2214;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C2A / ATCC 35395 / DSM 2834;
RX MEDLINE=21929760; PubMed=11932238;
RA Galagan J.E., Nusbaum C., Roy A., Endrizzi M.G., Macdonald P.,
RA Fitzhugh W., Calvo S., Engels R., Smirnov S., Atnoor D., Brown A.,
RA Allen N., Naylor J., Stange-Thomann N., DeArellano K., Johnson R.,
RA Linton L., McEwan P., McKernan K., Talamas J., Tirrell A., Ye W.,
RA Zimmer A., Barber R.D., Cann I., Graham D.E., Grahame D.A., Guss A.M.,
RA Hedderich R., Ingram-Smith C., Kuettnet H.C., Krzycki J.A.,
RA Leigh J.A., Li W., Liu J., Mukhopadhyay B., Reeve J.N., Smith K.,
RA Springer T.A., Unayam L.A., White O., White R.H., de Macario E.C.,
RA Perry J.G., Jarrell K.F., Jing H., Macario A.J.L., Paulsen I.,
RA Pritchett M., Sowers K.R., Swanson R.V., Zinder S.H., Lander E.,
RA Metcalf W.W., Birren B.;
RT "The genome of Methanosarcina acetivorans reveals extensive metabolic
RT and physiological diversity";
RL Genome Res. 12:532-542(2002).
DR EMBL: AE011107; AA007308.1; -.
DR InterPro: IPR003593; AAA_ATPase.
DR Pfam: PF00005; ABC_tran; 1.
DR ProDom: PD000006; ABC_transporter; 1.
DR SMART: SM00382; AAA; 1.
DR PROSITE: PS00211; ABC_TRANSPORTER; 1.
KW ATP-binding; Complete proteome.
SQ SEQUENCE 213 AA; 23676 MW; B548F93D94D109C9 CRC64;

  Query Match          50.6%; Score 42; DB 17; Length 213;
  Best Local Similarity 80.0%; Pred. No. 49;
  Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 GRELRRMSDE 13
   ||:|||||
Db 59 GRELRRMSDE 68

RESULT 14
Q8RCC4
ID Q8RCC4 PRELIMINARY; PRT; 260 AA.
AC Q8RCC4;
DT 01-JUN-2002 (TRENBLrel. 21, Created)
DT 01-JUN-2002 (TRENBLrel. 21, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE ABC-type amino acid periplasmic component.

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GN ARTI OR TTE0512.
OS Thermoanaerobacter tengcongensis.
OC Bacteria; Firmicutes; Clostridia; Thermoanaerobacteriales;
OC Thermoanaerobacteriaceae; Thermoanaerobacter.
OX NCBI_TaxID=119072;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MB4 / JCM 11007;
RX MEDLINE=21992816; PubMed=11997336;
RA Bao Q., Tian Y., Li W., Xu Z., Xuan Z., Hu S., Dong W., Yang J.,
RA Chen Y., Xue Y., Xu Y., Lai X., Huang L., Dong X., Ma Y., Ling L.,
RA Tan H., Chen R., Wang J., Yu J., Yang H.;
RT "A complete sequence of T. tengcongensis genome.";
RL Genome Res. 12:689-700(2002).
DR EMBL: AE013022; AA023788.1; -.
DR InterPro: IPR003439; ABC_transporter.
DR InterPro: IPR001311; SBP_glu_receptor.
DR InterPro: IPR001638; SBP_bac_3.
DR Pfam: PF00497; SBP_bac_3; 1.
DR SMART: SM00062; PBpb; 1.
DR PROSITE: PS00211; ABC_TRANSPORTER; 1.
DR PROSITE: PS01039; SBP_BACTERIAL_3; 1.
KW Complete proteome.
SQ SEQUENCE 260 AA; 28849 MW; F3D87ED16B18DCBC CRC64;

  Query Match          50.6%; Score 42; DB 16; Length 260;
  Best Local Similarity 58.3%; Pred. No. 61;
  Matches 7; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 5 RELRRMSDEFVD 16
   ||:|||||
Db 170 KOLNRVSDDEFMD 181

RESULT 15
Q8MZ90
ID Q8MZ90 PRELIMINARY; PRT; 297 AA.
AC Q8MZ90;
DT 01-OCT-2002 (TRENBLrel. 22, Created)
DT 01-OCT-2002 (TRENBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE AT26020P.
GN CG30456 OR CG15612 OR CG15613.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN [1]
RP SEQUENCE FROM N.A.
RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
RA Champe M., Chavez C., Dorsett V., Dresnek D., Farfan D., Frise B.,
RA George R., Gonzalez M., Guarin H., Kronmiller B., Li P., Liao G.,
RA Miranda A., Mungall C.J., Nunoo J., Pacleb J., Paragas V., Park S.,
RA Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,
RA Celniker S.;
RL Submitted (May-2002) to the EMBL/GenBank/DBSJ databases.
DR EMBL: AL113300; AA029305.1; -.
DR FlyBase: FBgn0050456; CG30456.
DR InterPro: IPR000219; RhogEF.
DR Pfam: PF00621; RhogEF; 1.
DR PROSITE: PS0010; DH_2; 1.
SQ SEQUENCE 297 AA; 35804 MW; 58FDBF3DF688D99A CRC64;

  Query Match          50.8%; Score 42; DB 5; Length 297;
  Best Local Similarity 53.3%; Pred. No. 70;
  Matches 8; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFV 15
   ||:|||||
Db 3 QNRNEDLRKLFDEFL 17

```

Search completed: September 15, 2003, 17:25:49
Job time : 18.3714 secs

1

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 15, 2003, 17:18:16 ; Search time 7.2 Seconds
(without alignments)
213.708 Million cell updates/sec

Title: US-09-544-664-29

Perfect score: 83

Sequence: 1 QRYGRELRRMSDEFVD 16

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR_76.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Score	Match	Length	DB ID	Description
1	73	88.0	204	2 A55671	bad protein - mous
2	45	54.2	564	2 H75403	glycosyl hydrolase
3	44	53.0	1248	2 G83278	cobalamin biosynth
4	42	50.6	876	2 E89949	valine-trna ligase
5	41.5	50.0	191	2 AD3414	hypothetical cytos
6	41	49.4	447	2 F83356	hypothetical prote
7	41	49.4	577	2 T40297	membrane transport
8	41	49.4	858	2 A44919	GCR3 protein - yea
9	40	48.2	84	2 F84388	hypothetical prote
10	40	48.2	109	2 E95043	conserved hypotet
11	40	48.2	113	2 D97913	conserved hypotet
12	40	48.2	219	2 A75088	hypothetical prote
13	40	48.2	275	2 E91102	probable enzyme [i
14	40	48.2	275	2 A85948	probable enzyme yg
15	40	48.2	275	2 F63076	hypothetical prote
16	40	48.2	335	2 T52577	gibberellin 2beta-
17	40	48.2	360	2 B83311	hypothetical prote
18	40	48.2	411	2 F87644	transcription regu
19	40	48.2	429	2 JCA986	site-specific DNA-
20	40	48.2	445	2 G97123	probable Fe-S oxid
21	40	48.2	631	2 G70188	transcription init
22	40	48.2	5138	2 B96695	hypothetical prote
23	39.5	47.6	198	2 E87441	conserved hypotet
24	39.5	47.6	414	2 B84275	hypothetical prote
25	39	47.0	73	2 AC3365	hypothetical prote
26	39	47.0	172	2 B71339	probable cationic
27	39	47.0	207	2 B95348	hypothetical prote
28	39	47.0	220	2 F72289	oxidoreductase, so
29	39	47.0	275	2 C69808	transporter homolo

```

30      39      47.0      331      2      E90121      DNA repair protein
31      39      47.0      360      2      D86200      protein FlzKil1.20
32      39      47.0      365      2      S42107      RAD51 protein homo
33      39      47.0      380      2      T32163      hypothetical prote
34      39      47.0      383      2      T31738      hypothetical prote
35      39      47.0      418      1      FOXRL2      sigma 2 protein -
36      39      47.0      418      1      FOXB3D      sigma 2 protein -
37      39      47.0      432      2      AB0558      trigger factor [im
38      39      47.0      503      1      CTBPRH      site-specific DNA-
39      39      47.0      536      2      F90299      acylaminoacyl-pept
40      39      47.0      689      2      T29772      hypothetical prote
41      39      47.0      880      1      SVBSVS      valine-trna ligase
42      39      47.0      959      1      B71405      probable kinesin -
43      39      47.0      985      2      AE2452      two-component hybr
44      39      47.0      1805      2      T02712      similar to late em
45      39      47.0      1967      2      S64604      hypothetical prote

```

ALIGNMENTS

RESULT 1

A55671

bad protein - mouse

C:Species: Mus musculus (house mouse)

C>Date: 03-Mar-1995 #sequence_revision 03-Mar-1995 #text_change 05-Nov-1999

C:Accession: A55671

R:Yang, E.; Zha, J.; Jockel, J.; Boise, L.H.; Thompson, C.B.; Korsmeyer, S.J.

Cell 80, 285-291, 1995

A:Title: Bad, a heterodimeric partner for Bcl-x-L and Bcl-2, displaces Bax and promot

A:Reference number: A55671; MUID:95136361; PMID:7834748

A:Accession: A55671

A>Status: Preliminary; not compared with conceptual translation

A:Molecule type: mRNA

A:Residues: 1-204 <YAN>

A:Cross-references: GB:L37296; NID:9639778; PIDN:AAA64465.1; PID:9639779

C:Keywords: heterodimer

Query Match 88.0%; Score 73; DB 2; Length 204;

Best Local Similarity 100.0%; Pred. No. 0.00011;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

Qy      1 QRYGRELRRMSDEF 14
        |||||
Db      145 QRYGRELRRMSDEF 158

```

RESULT 2

H75403

glycosyl hydrolase, family 13 - Deinococcus radiodurans (strain R1)

C:Species: Deinococcus radiodurans

C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 17-Mar-2000

C:Accession: H75403

R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.

M.; Shen, M.O.; Vanathavan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zalewski, C.

S.; Smith, H.O.; Veneter, J.C.; Fraser, C.M.

Science 286, 1571-1577, 1999

A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.

A:Reference number: A75250; MUID:20036896; PMID:10567266

A:Accession: H75403

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-564 <WHI>

A:Cross-references: GB:AE001993; GB:AE000513; NID:96459123; PIDN:AAF10944.1; PID:964;

A:Experimental source: strain R1

C:Genetics:

A:Gene: DR1375

A:Map position: 1

C:Superfamily: alpha-glucosidase; alpha-amylase core homology

Query Match 54.2%; Score 45; DB 2; Length 564;

Best Local Similarity 64.3%; Pred. No. 14;

Matches 9; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

```

C;Accession: AD3414
R;DelVecchio, V.G.; Kapetral, V.; Redkar, R.J.; Patra, G.; Mijer, C.; Los, T.; Ivanov
.: Mazur, M.; Goltzman, E.; Selkov, E.; Elzer, P.H.; Hagius, S.; O'Callaghan, D.; Let
Proc Natl Acad Sci U S A. 99, 443-448, 2002
A;Title: The genome sequence of the facultative intracellular pathogen Brucella melit
A;Reference number: AD3252; PMID:11756688
A;Accession: AD3414
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-191 <KUR>
A;Cross-references: GB:AE008917; PIDN:AAL52479.1; PID:g17983287; GSPDB:GN00190
A;Experimental source: strain 16M
C;Genetics:
A;Gene: BME11298
A;Map position: 1

Query Match 50.0%; Score 41.5; DB 2; Length 191;
Best Local Similarity 33.3%; Pred. No. 18;
Matches 11; Conservative 2; Mismatches 3; Indels 17; Gaps 1;

Qy 1 QRYGR-----ELFRMSDEFVD 16
Db 131 QKGRKVVSVTITTOPAMISDELRRQADHFID 163

RESULT 6
F85356
hypothetical protein AT4g30490 [imported] - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 16-Feb-2001
C;Accession: F85356
R;Anonymous, The European Union Arabidopsis Genome Sequencing Consortium, The Cold Sp
Nature 402, 769-777, 1999
A;Title: Sequence and analysis of chromosome 4 of the plant Arabidopsis thaliana.
A;Reference number: A85001; MUID:20083488; PMID:10617198
A;Accession: F85356
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-447 <STO>
A;Cross-references: GB:NC_001268; NID:g7269950; PIDN:CAB79767.1; GSPDB:GN00140
C;Genetics:
A;Gene: AT4g30490
A;Map position: 4

Query Match 49.4%; Score 41; DB 2; Length 447;
Best Local Similarity 56.7%; Pred. No. 52;
Matches 8; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 5 RELRRMSDEFVD 16
Db 84 RELQRLVDELVD 95

RESULT 7
T40297
membrane transporter - fission yeast (Schizosaccharomyces pombe)
C;Species: Schizosaccharomyces pombe
C;Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 18-Feb-2000
C;Accession: T40297
R;Lyne, M.; Wood, V.; Rajandream, M.A.; Barrell, B.G.; Hilbert, H.; Moestl, D.; Duest
submitted to the EMBL Data Library, May 1998
A;Reference number: #21919
A;Accession: T40297
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-577 <LYN>
A;Cross-references: EMBL:AL023589; PIDN:CAA19050.1; GSPDB:GN00067; SPDB:SPEC36.02c
A;Experimental source: strain 972h-; cosmid c36
C;Genetics:
A;Gene: SPDB:SPEC36.02c
A;Map position: 2
C;Superfamily: benomyl/methotrexate resistance protein

Cobalamin biosynthetic protein CbN PA2944 [imported] - Pseudomonas aeruginosa (strain F
C;Species: Pseudomonas aeruginosa
C;Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 02-Mar-2001
C;Accession: G83278
R;Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.; Bu
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Harbig, K.; Lim,
.: Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A;Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic patho
A;Reference number: A82950; MUID:20437337; PMID:10984043
A;Accession: G83278
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-1248 <SPO>
A;Cross-references: GB:AE004720; GB:AE004091; NID:g9949032; PIDN:AAG06332.1; GSPDB:GN001
A;Experimental source: strain PA01
C;Genetics:
A;Gene: cbN; PA2944
C;Superfamily: Rhodobacter capsulatus magnesium-protoporphyrin O-methyltransferase

Query Match 53.0%; Score 44; DB 2; Length 1248;
Best Local Similarity 50.0%; Pred. No. 47;
Matches 9; Conservative 4; Mismatches 3; Indels 2; Gaps 1;

Qy 1 QRYG--RELRRMSDEFVD 16
Db 615 ESYGPLRLERLADEFVD 632

RESULT 4
E89949
valine-tRNA ligase [imported] - Staphylococcus aureus (strain N315)
C;Species: Staphylococcus aureus
C;Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 22-Oct-2001
C;Accession: E89949
R;Kuroda, M.; Ohta, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; Oguc
ma, A.; Mizutani-Ui, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K.;
C.; Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramatsu, K.
Lancet 357, 1225-1240, 2001
A;Title: Whole genome sequencing of methicillin-resistant Staphylococcus aureus.
A;Reference number: A89758; MUID:21311952; PMID:11418146
A;Accession: E89949
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-876 <KUR>
A;Cross-references: GB:BA000018; PID:g13701460; PIDN:BA842754.1; GSPDB:GN00149
A;Experimental source: strain N315
C;Genetics:
A;Gene: valS
C;Superfamily: valine-tRNA ligase

Query Match 50.6%; Score 42; DB 2; Length 876;
Best Local Similarity 61.5%; Pred. No. 70;
Matches 8; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 4 GRELRMSDEFVD 16
Db 251 GRELPILADEYVD 263

RESULT 5
AD3414
hypothetical cytosolic protein BME11298 [imported] - Brucella melitensis (strain 16M)
C;Species: Brucella melitensis
C;Date: 01-Feb-2002 #sequence_revision 01-Feb-2002 #text_change 01-Feb-2002

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Query Match          49.4%; Score 41; DB 2; Length 577;
Best Local Similarity 57.1%; Pred. No. 67;
Matches 8; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 YGRELRMSDEFVD 16
    :||:|||||
Db 563 FGRKIRMSKMAVD 576

RESULT 8
A44919
GCR3 protein - yeast (Saccharomyces cerevisiae)
N:Alternate names: protein YMR564.07; protein YMR125W
C:Species: Saccharomyces cerevisiae
C:Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 29-Oct-1999
C:Accession: A44919; S53055; S54494
R:Uemura, H.; Jigami, Y.
J. Bacteriol. 174, 5526-5532, 1992
A:Title: GCR3 encodes an acidic protein that is required for expression of glycolytic genes
A:Reference number: A44919; MUID:92380925; PMID:1512188
A:Accession: A44919
A:Molecule type: DNA
A:Residues: 1-858 <UDEM>
A:Cross-references: GB:D10224; NID:G464221; PIDN:BAA01076.1; PID:d1001545; PID:G464222
A:Note: sequence extracted from NCBI backbone (NCBIN:112104, NCBIP:112106)
R:Badcock, K.; Churcher, C.
Submitted to the EMBL Data Library, March 1995
A:Reference number: S53055
A:Accession: S53055
A:Molecule type: DNA
A:Residues: 339-858 <BAD>
A:Cross-references: EMBL:Z48622; NID:G728663; PIDN:CAR88550.1; PID:G728664; MIPS:YMR125W
R:Lye, G.; Churcher, C.M.
Submitted to the EMBL Data Library, May 1995
A:Reference number: S54014
A:Accession: S54494
A:Molecule type: DNA
A:Residues: 'MFNRKRG', 6-499 <LYE>
A:Cross-references: EMBL:Z49273; NID:G809577; PIDN:CAR89274.1; PID:G809584; MIPS:YMR125W
C:Genetics:
A:Gene: SGD:STO1; GCR3
A:Cross-references: MIPS:YMR125W; SGD:S0004732
A:Map position: 13R
C:Keywords: DNA binding; nucleus

Query Match          49.4%; Score 41; DB 2; Length 858;
Best Local Similarity 40.0%; Pred. No. 1e+02;
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 1 QRYGRELRRMSDEFV 15
    :||:|||||
Db 818 RYISHEYRELADRFI 832

RESULT 9
F84388
hypothetical protein Vng2379h [imported] - Halobacterium sp. NRC-1
C:Species: Halobacterium sp. NRC-1
C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C:Accession: F84388
R:Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.;
Leithausen, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jablo
Jung, K.H.; Alam, M.; Freitas, I.
Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000
A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li
A:Title: Genome sequence of Halobacterium species NRC-1.
A:Reference number: A84160; MUID:20504483; PMID:11016950
A:Accession: F84388
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-84 <STO>
A:Cross-references: GB:AF004437; NID:G10581786; PIDN:AAG20474.1; GSPDB:GN00138
C:Genetics:

A:Gene: VNG2379H

Query Match          48.2%; Score 40; DB 2; Length 84;
Best Local Similarity 66.7%; Pred. No. 14;
Matches 8; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 RYGRELRMSDE 13
    ||| | |||
Db 66 RYGTASMRDE 77

RESULT 10
B95043
conserved hypothetical protein SP0372 [imported] - Streptococcus pneumoniae (strain ;
C:Species: Streptococcus pneumoniae
C:Date: 03-Aug-2001 #sequence_revision 03-Aug-2001 #text_change 03-Aug-2001
C:Accession: B95043
R:Tettelin, H.; Nelson, K.E.; Paulsen, I.T.; Eisen, J.A.; Read, T.D.; Peterson, S.; I
on, J.D.; Umayam, L.A.; White, O.; Salzberg, S.L.; Lewis, M.R.; Radune, D.; Holtzapp
son, R.; Hickey, E.K.; Holt, I.E.
Science 293, 498-506, 2001
A:Authors: Loftus, B.J.; Yang, F.; Smith, H.O.; Venter, J.C.; Dougherty, B.A.; Morris
A:Title: Complete Genome Sequence of a virulent isolate of Streptococcus pneumoniae.
A:Reference number: A95000; MUID:21357209; PMID:11463916
A:Accession: B95043
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-109 <KUR>
A:Cross-references: GB:AE005672; PIDN:AAK74539.1; PID:g14971841; GSPDB:GN00164; TIGR
A:Experimental source: strain TIGR4
C:Genetics:
A:Gene: SP0372

Query Match          48.2%; Score 40; DB 2; Length 109;
Best Local Similarity 45.0%; Pred. No. 18;
Matches 9; Conservative 4; Mismatches 3; Indels 4; Gaps 1;

QY 1 QRYGRELRRMS----DEFVD 16
    | :|||: | :|||:
Db 14 QEFGEVGRYKNKVEVDFLD 33

RESULT 11
D97913
conserved hypothetical protein spr0332 [imported] - Streptococcus pneumoniae (strain
C:Species: Streptococcus pneumoniae
C:Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 22-Oct-2001
C:Accession: D97913
R:Hoskins, J.A.; Alborn Jr., W.; Arnold, J.; Blaszcak, L.; Burgett, S.; DeGoff, B.S
e, R.; Leblanc, D.J.; Lee, L.N.; Lefkowitz, E.J.; Lu, J.; Matsushima, P.; McAhren, S
Y, P.; Sun, P.M.; Winkler, M.E.
J. Bacteriol. 183, 5709-5717, 2001
A:Authors: Yang, Y.; Young-Bellido, M.; Zhao, G.; Zook, C.; Baltz, R.H.; Jaskunas, S
A:Title: Genome of the Bacterium Streptococcus pneumoniae Strain R6.
A:Reference number: A97872; MUID:21429245; PMID:11544234
A:Accession: D97913
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-113 <KUR>
A:Cross-references: GB:AE007317; PIDN:AAK99136.1; PID:g15457889; GSPDB:GN00174
C:Genetics:
A:Gene: spr0332

Query Match          48.2%; Score 40; DB 2; Length 113;
Best Local Similarity 45.0%; Pred. No. 19;
Matches 9; Conservative 4; Mismatches 3; Indels 4; Gaps 1;

QY 1 QRYGRELRRMS----DEFVD 16
    | :|||: | :|||:
Db 18 QEFGEVGRYKNKVEVDFLD 37

RESULT 12

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A75088
Hypothetical protein PAB1640 - Pyrococcus abyssi (strain Orsay)
C:Species: Pyrococcus abyssi
C:Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999
C:Accession: A75088
R:anonymous, Genoscope
submitted to the EMBL Data Library, July 1999
A:Description: Pyrococcus abyssi genome sequence: insights into archaeal chromosome stru
A:Reference number: A75001
A:Accession: A75088
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-219 <NAN>
A:Cross-references: GB:A749286; GB:AL096836; NID:g5458366; PIDN:CAB50006.1; PID:e151590
A:Experimental source: strain Orsay
C:Genetics:
A:Gene: PAB1640

Query Match      48.2%; Score 40; DB 2; Length 219;
Best Local Similarity 47.1%; Pred. No. 37;
Matches 8; Conservative 4; Mismatches 3; Indels 2; Gaps 1;

QY      2  YGRELRRMS--DEFVD 16
      ||| | :| :| :| :|
Db      104  YGNEFQVSPQENFID 120

RESULT 13
E91102
probable enzyme [imported] - Escherichia coli (strain O157:H7, substrain RIMD 0509952)
C:Species: Escherichia coli
C:Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001
C:Accession: E91102
R:Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.-G.;
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and gene
A:Reference number: A99629; MUID:21156231; PMID:11258796
A:Accession: E91102
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-275 <HAY>
A:Cross-references: GB:BA000007; PIDN:BA037212.1; PID:g13363261; GSPDB:GN00154
A:Experimental source: strain O157:H7, substrain RIMD 0509952
C:Genetics:
A:Gene: ECs3789
C:Superfamily: naphthoate synthase; enoyl-CoA hydratase homology

Query Match      48.2%; Score 40; DB 2; Length 275;
Best Local Similarity 50.0%; Pred. No. 46;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY      3  YGRELRRMSDEFVD 16
      ||| | :| :| :| :|
Db      35  YGKLNALSKVFID 48

RESULT 14
A85948
probable enzyme yfg [imported] - Escherichia coli (strain O157:H7, substrain EDL933)
C:Species: Escherichia coli
C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001
C:Accession: A85948
R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
iller, L.; Grobeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A:Reference number: A85480; MUID:21074935; PMID:11206551
A:Accession: A85948
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-275 <STO>
A:Cross-references: GB:AE005174; NID:g12517451; PIDN:AA058045.1; GSPDB:GN00145; UWGP:242

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A:Experimental source: strain O157:H7, substrain EDL933
C:Genetics:
A:Gene: yfg
C:Superfamily: naphthoate synthase; enoyl-CoA hydratase homology

Query Match      48.2%; Score 40; DB 2; Length 275;
Best Local Similarity 50.0%; Pred. No. 46;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY      3  YGRELRRMSDEFVD 16
      ||| | :| :| :| :|
Db      35  YGKLNALSKVFID 48

RESULT 15
F65076
hypothetical protein b2919 - Escherichia coli (strain K-12)
C:Species: Escherichia coli
C:Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 01-Mar-2002
C:Accession: F65076
R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.;
.A.; Rose, D.J.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A:Title: The complete genome sequence of Escherichia coli K-12.
A:Reference number: A64720; MUID:97426617; PMID:9278503
A:Accession: F65076
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-275 <BLAT>
A:Cross-references: GB:AF000375; GB:U00096; NID:g1789282; PIDN:AAC75956.1; PID:g1789;
A:Experimental source: strain K-12, substrain M01655
C:Superfamily: naphthoate synthase; enoyl-CoA hydratase homology
F:40-192/Domain: enoyl-CoA hydratase homology <ECH>

Query Match      48.2%; Score 40; DB 2; Length 275;
Best Local Similarity 50.0%; Pred. No. 46;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY      3  YGRELRRMSDEFVD 16
      ||| | :| :| :| :|
Db      35  YGKLNALSKVFID 48

Search completed: September 15, 2003, 17:27:03
Job time : 9.2 secs

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